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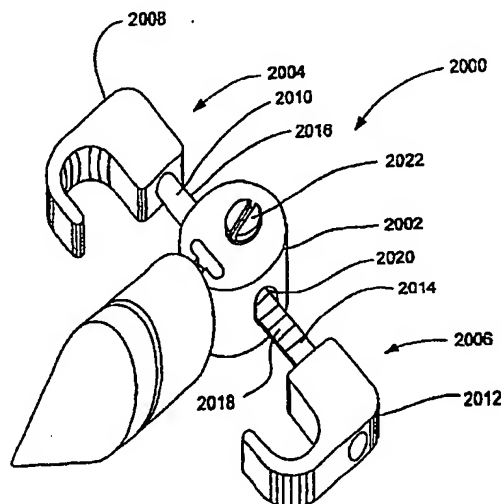
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(54) Title: SUPPLEMENTAL SPINE FIXATION DEVICE AND METHOD



(57) Abstract: A supplemental spine fixation device and method is used in association with a primary fixation device. The supplemental spine fixation device includes a guide (2026) and spacer (2041) for distracting apart adjacent spinous processes and the device has hook members (2008, 2012) which hook about the first and second spinous processes. With the spinous processes distracted and the hook members about the spinous processes, the hook members can be rigidly secured to a hub (2002) in order to rigidly affix the spinous processes about the spacer. With such an arrangement, additional force is placed on vertebral bodies in order to encourage fusion. In addition, the rigidity between the spinous processes assures that the vertebral bodies will be held rigidly in place in order to promote bone growth and fusion.

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SUPPLEMENTAL SPINE FIXATION DEVICE AND METHOD

Cross-References to Related Applications:

5 This application is a continuation-in-part of U.S. Patent Application No. 09/473,173 filed on December 28, 1999 and entitled SPINE DISTRACTION IMPLANT (which is a continuation of U.S. Patent Application No. 09/179,570 filed on October 27, 1998 and entitled SPINE DISTRACTION IMPLANT, now U.S. Patent No. 6,048,342 issued April 11, 2000), which is a continuation-in-part of U.S. Patent Application No. 10 09/474,037 filed on December 28, 1999 and entitled SPINE DISTRACTION IMPLANT (which is a continuation of U.S. Patent Application No. 09/175,645 filed on October 20, 1998 and entitled SPINE DISTRACTION IMPLANT, now U.S. Patent No. 6,068,630 issued May 30, 2000. All of the above applications and patents are 15 incorporated herein by reference.

Field of the Invention:

20 The present invention is directed to supplemental spine fixation devices and methods which are used as an adjunct to a primary spine fusion device, such as by way of example only, an interbody fusion device.

Background of the Invention:

25 A common procedure for handling pain associated with degenerative spinal disk disease is the use of devices for fusing together two or more adjacent vertebral bodies. The procedure is known by a number of terms, one of which is interbody fusion. Interbody fusion can be accomplished through the use of a number of

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devices and methods known in the art. These include screw arrangements, solid bone implant methodologies, and fusion devices which include a cage or other mechanism which is packed with bone and/or bone growth inducing substances. All of the above are implanted
5 between adjacent vertebral bodies in order to fuse the vertebral bodies together, alleviating associated pain.

Associated with such primary fusion devices and methods are supplemental devices which assist in the fusion process. These supplemental devices assist during the several month period when bone
10 from the adjacent vertebral bodies is growing together through the primary fusion device in order to fuse the adjacent vertebral bodies. During this period it is advantageous to have the vertebral bodies held immobile with respect to each other so that sufficient bone growth can be established.

15 Such supplemental devices can include hook and rod arrangements, screw arrangements, and a number of other devices which include straps, wires, and bands, all of which are used to immobilize one portion of the spine relative to another.

All of these devices generally require extensive surgical
20 procedures in addition to the extensive procedure surrounding the primary fusion implant.

It would be advantageous if the device and procedure for supplemental spine fixation were as simple and easy to perform as possible, and would leave intact all bone, ligament, and other tissue
25 which comprise and surround the spine.

Accordingly, there needs to be developed procedures and implants which are minimally invasive and are supplemental to spine fixation devices and methods.

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Summary of the Invention:

The present invention is directed to providing a minimally invasive supplemental spine fixation implant and method for alleviating discomfort associated with the spine.

5 The present invention provides for a method and apparatus for assisting in the fusing together of vertebral bodies of the spine. One of the features and purposes of the invention is to immobilize the vertebral bodies while spine fusion is accomplished. Generally fusion requires upwards of six months for bone cells from the upper and lower vertebral
10 bodies to grow towards each other, generally through a primary fusion device.

 In order to assist in the fusing process, the supplemental spinal fixation device and method of the invention immobilizes the vertebral bodies by immobilizing the respective spinous processes extending
15 therefrom. In addition, the present invention and method can be used to distract apart the posterior sides of the vertebral bodies in order to put additional force and compression on the anterior sides of the vertebral bodies, further assisting in the interbody fusion process.

 The present invention and method is minimally invasive such that
20 it does not add to the trauma of the primary fusion procedure, especially if the fusion procedure is from a posterior approach. With an anterior fusion approach additional posterior incisions are required. However, these are minimal when compared to other devices and methods.

 Accordingly an object of the present invention is to increase the
25 rigidity and stability with respect to the adjacent spinous process and vertebral bodies in order to promote interbody fusion between the vertebral bodies.

 It is further an object of the present invention to be as minimally invasive as possible.

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It is yet a further object of the present invention to provide for an implant and method which does not require modification of the bone, ligaments, or adjoining tissues. In other words, it is an object of the present invention to provide for an implant and method which does not
5 require that the bone be reshaped, notched, or in any way modified. Further it is an object of the present invention to provide for an implant and method which does not require that any of the ligaments associated with the spinous processes be altered.

It is a further object of the present invention to provide for an
10 implant and method which can be inserted from one side of adjacent spinous processes, in order to immobilize the spinous processes and resultingly immobilize the adjacent vertebral bodies. By addressing the spinous processes from one side, the objects and advantages of a minimally invasive procedure, with reduced trauma, can be
15 accomplished.

It is another object of the present invention to provide for a device and method which provides for distraction of the spinous processes in order to place pressure on at least the anterior portion of the vertebral bodies in order to assist in the primary fusion.

20 It is still a further object of the present invention to provide for an implant and method which can increase the space between spinous processes in order to adjust the height between vertebral bodies.

It is yet a further object of the present invention to provide for a device which has securing and/or hook elements which can easily and
25 conveniently be secured about the spinous processes, which hook devices are preferably designed in order to accommodate the shape of the spinous processes and are preferably swivelable or pivotable in order to accommodate the position and shape of one spinous processes relative to another.

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It is another object of the invention to provide for a device which has several degrees of freedom in order to allow a portion of the device to be positioned between spinous processes in order to distract apart the spinous processes and other portions of the device to engage the spinous processes in order to rigidly immobilize the spinous processes. These degrees of freedom allow the device to conform to the bones, ligaments, and tissues of each individual patient. Thus, the present device allows for adjustments along two and three axes in order to successfully distract and immobilize spinous processes.

It is yet a further object of the present invention to have at least one portion of the device selectably positionable with respect to other portions of the device in order to accommodate the anatomy of the spine and in particular of the spinous processes.

It is still a further object of the present invention to provide for a device and method which can be used with both primary anterior or posterior interbody fusion.

Accordingly, it is an object and aspect of the invention to provide a device and method for augmentation of single or multiple level lumbar spinal fusion. Ideally the fusion and the device and method of this invention are addressed at the L4/L5 vertebral bodies and above, and also at the L5/S1 vertebral bodies. The device and method can also be used with other vertebral bodies located along the spine.

The present invention provides for rigidity without risk to the neural elements. The present invention is cost effective and minimally invasive.

Accordingly, an aspect of the present invention includes an implant for rigidly positioning spinous processes, which implant includes a first means adapted for engaging the first spinous process and a second means adapted for engaging the second spinous process. The

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implant includes a body means adapted for positioning between the first spinous process and the second spinous process and a hub means for engaging the first means, the second means, and the body means.

Further, the invention includes at least one of the hub means and
5 the body means allowing for the body means to move relative to at least one of the first and second means.

In a further aspect and object of the present invention, an implant includes a first hook adapted to engage a first spinous process and a second hook adapted to engage a second spinous process. The
10 implant has a body adapted to the position between the spinous processes and a hub to which mounts the first and second hooks and the body. The body is moveable relative to at least one of the first and second hooks.

It is further an aspect and object of the present invention to
15 provide an implant for rigidly positioning spinous processes as an adjunct to spine fusion, where the improvement includes a sleeve position between adjacent spinous processes.

It is a further aspect of the present invention to provide an implant for rigidly positioning spinous processes as an adjunct to spine fusion
20 wherein the improvement comprises a sleeve or spacer positioned between adjacent spinous processes and a first hook which is adapted to engage a first spinous process and a second hook which is adapted to engage a second spinous process.

The method of the present invention is for rigidly positioning a first
25 spinous process relative to a second spinous process and includes the steps in any desired order of placing a first hook around a first spinous process and a second hook around a second spinous process. The steps include placing a sleeve or spacer between the first and second

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spinous processes, which spacer mounts to a hub. The hub is used to interlock the first hook relative to the second hook.

5 Other embodiments of the implants and methods, within the spirit and scope of the invention, can be used to increase the volume of the spinal canal thereby alleviating restrictions on vessels and nerves associated therewith, and accordingly thereby alleviating pain.

Brief Description of the Figures

10 Figs. 1 and 2 depict an embodiment of an implant of the invention which is adjustable in order to select the amount of distraction required. Fig. 1 depicts the implant in a more extended configuration than does Fig. 2.

15 Figs. 3a and 3b depict side and end views of a first forked and of the embodiment of Fig. 1.

Figs. 4a and 4b depict side sectioned and end views of an interbody piece of the implant of Fig. 1.

Figs. 5a and 5b depict side and end views of a second forked end of the embodiment of Fig. 1.

20 Figs. 6, 7, 8, 9 and 10 depict apparatus and method for another embodiment of the present invention for creating distraction between adjacent spinous processes.

Figs. 11, 12 and 13 depict yet a further embodiment of the invention for creating distraction between adjacent spinous processes.

25 Figs. 14 and 15 depict a further apparatus and method of an embodiment of the invention for creating distraction.

Figs. 16, 16a, and 17 depict yet another embodiment of the present invention.

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Figs. 18, 19 and 20 depict yet a further apparatus and method of the present embodiment.

Figs. 21 and 22 depict still a further embodiment of the present invention.

5 Figs. 23, 24 and 25 depict another embodiment of the present invention.

Figs. 26, 27 and 28 depict another embodiment of the invention.

Figs. 29 and 30 depict side elevational views of differently shaped implants of embodiments of the present invention.

10 Figs. 31, 32 and 33 depict various implant positions of an apparatus of the present invention.

Figs. 34 and 35 depict yet another apparatus and method of the present invention.

15 Figs. 36, 37 and 38 depict three different embodiments of the present invention.

Figs. 39 and 40 depict yet another apparatus and method of an embodiment of the present invention.

Figs. 41, 42 and 43 depict yet further embodiments of an apparatus and method of the present invention.

20 Fig. 44 is still a further embodiment of an implant of the invention.

Fig. 45 is yet another depiction of an apparatus and method of the invention.

Figs. 46 and 47 depict still a further apparatus and method of an embodiment of the invention.

25 Figs. 48, 49, 50 and 51 depict yet a further apparatus and method of the invention.

Figs. 52, 53, 54, 55a and 55b depict another apparatus and method of the invention.

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Figs. 56, 57 and 58 depict yet a further apparatus and method of the invention.

Figs. 59 and 60 depict still a further embodiment of the invention.

Fig. 61 depict another embodiment of the invention.

5 Figs. 62 and 63 depict yet another embodiment of the present invention.

Figs. 64 and 65 depict still a further embodiment of the present invention.

Fig. 66 depicts another embodiment of the invention.

10 Figs. 67 and 68 depict yet another embodiment of the present invention.

Figs. 69, 70, 71 and 71a depict a further embodiment of the present invention.

Figs. 72 and 73 depict still another embodiment of the invention.

15 Figs. 74, 75, 76, 77, and 78 depict still other embodiments of the invention.

Figs. 79, 80, 80a, 81, 82, 83, 83a, 84, 85, 86 and 87 depict still a further embodiment of the present invention.

20 Figs. 88, 89, 90 and 91 depict yet another embodiment of the present invention.

Figs. 92, 92a, 92b, 93, 93a, 93b, 93c, 93d, 94, 94a, 94b, 95, 95a, and 96, depict still a further embodiment of the present invention wherein a sleeve is provided which is capable of deflecting response to relative motion between the spinous processes.

25 Fig. 97 depicts still another embodiment of the present invention.

Fig. 98 depicts yet a further embodiment of the present invention.

Figs. 99 and 100 depict still another embodiment of the present invention including an insertion tool.

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Figs. 101, 102, 102a, 103, 104, 105, 106, and 107 depict still a further embodiment of the present invention.

Figs. 108, 109, and 110 depict still another embodiment of the present invention.

5 Figs. 111, 112, 113, 114, 115, 116, and 117 depict yet another embodiment of the present invention.

Fig. 118 depicts a graph showing characteristics of a preferred material usable with several of the embodiments of the present invention.

10 Figs. 119a and 119b depict side and plan views of still a further embodiment of the present invention.

Figs. 120a and 120b depict side and plan views of the second wing which can be used in conjunction with the embodiment of the invention of Figs. 119a and 119b.

15 Figs. 121a and 121b depict side and plan views of the first wing and central body of the embodiment of the invention depicted in Figs. 119a and 119b.

Figs. 122a, 122b, and 122c depict top, side and end views of a guide which is a portion of the embodiment of the invention of Figs. 119a and 119b.

20 Figs. 123a and 123b depict an end view and a cross-sectioned view respectfully of the sleeve of the embodiment of the invention of Figs. 119a and 119b.

Figs. 124a, 124b and 124c depict a view of the embodiment of the invention of Figs. 119a and 119b taken through line 124-124 in Fig. 119b shown in with the sleeve in various positions relative to a first wing.

25 Fig. 125 depicts an alternative embodiment of the invention as depicted in Figs. 119a and 119b.

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Fig. 126 depicts yet a further alternative embodiment of the invention depicted in Figs. 119a and 119b.

Fig. 127 depicts yet a further embodiment of the invention as depicted in Figs. 119a and 119b.

5 Fig. 128 is still a further embodiment of the invention as depicted in Fig. 93a.

Fig. 129 depicts still a further embodiment of the invention as depicted in Figs. 119a and 119b.

10 Fig. 130 is a perspective view of a first embodiment of the invention.

Fig. 131 is an exploded view of the embodiment of the invention of Fig. 130. Fig. 131a and 131b are alternative components of the embodiment of Fig. 131.

15 Fig. 132 is a plan view of the embodiment of the invention of Fig. 130.

Figs. 133a, 133b, 133c, and 133d are perspective, first end, second end, and sectional views of a spacer or sleeve of the embodiment of the invention depicted in Fig. 130.

20 Fig. 134 is a cross sectional view of an embodiment of the invention taken through line 134-134 in Fig. 132.

Figs. 135a-135f are various views of an embodiment of the hook mechanism of the embodiment of the invention of Fig. 130.

Fig. 136 is a schematical representation of an embodiment of the invention as positioned with respect to adjacent spinous processes.

25 Fig. 137 is a perspective view of another embodiment of the invention.

Fig. 138 is an exploded view of the embodiment of the invention of Fig. 137.

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Fig. 138a is an alternative component of the embodiment of Fig. 137.

Fig. 138b is an upside down perspective view of a component of the embodiment of Fig. 138.

5 Fig. 139 is a plan view of the embodiment of the invention of Fig. 137.

Fig. 140 is a partial section view taken through line 140-140 of Fig. 139.

10 Fig. 141 is an exploded view of yet another embodiment of the invention. Fig. 141a is an upside down perspective view of a component of the embodiment of Fig. 141.

Fig. 142 is a sectional view of a body portion of the embodiment of the invention of Fig. 141 taken through line 142-142.

Fig. 143 is a top view of the body portion shown in Fig. 142.

15 Fig. 144 is a sectional view of yet another embodiment of a body portion of the invention.

Fig. 145 is a perspective view of yet a further embodiment of the body portion of the invention.

20 Figs. 146a, 146b, and 146c depict yet a further embodiment of a body portion of the invention.

Figs. 147a and 147b are side and top views of yet another embodiment of the invention depicting a mechanism for adjusting the positions of the hook mechanisms of, for example, the embodiment of the invention of Figs. 130, 137, and 141.

25 Figs. 148a and 148b are sectional top and side views of yet another embodiment of the invention for adjusting the position of the hook mechanisms.

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Figs. 149a and 149b are perspective and side views of yet a further mechanism of an embodiment of the invention for adjusting the position of hook mechanisms of the invention.

5 Fig. 150 is a perspective view of yet a further embodiment of the invention.

Fig. 151 is a perspective view of an embodiment of the invention which is addressable to multiple levels of spinous processes.

Detailed Description of the Preferred Embodiment

10 The present invention, although directed to embodiments for providing supplemental spine fixation devices and methodologies depicted in Figs. 130 to 151, have some of the same functionalities, features, design characteristics, and materials as previously described in the embodiments depicted and described in Figs. 1 to 129. Figs. 1
15 to 129 are directed to spine distraction implant and method used in distracting apart spinous processes in order to relieve pain associated with the spine such as, by way of example only, the pain associated with spinal stenosis. Accordingly, as appropriate, and even if not specifically mentioned in each inventive description of Figs. 130 to 151, many of the
20 design characteristics, features, functionalities, materials, measurements, dimensions, purposes, aspects, and objects of the devices in Figs. 1 to 129 are applicable to the present invention.

Embodiment of Figs. 1-5a, 5b

25 A first embodiment of the invention is shown in Figs. 1-5a, 5b. Implant 20 includes first and second forked ends 22 and 24, each defining a saddle 26, 28 respectively. The forked ends 22, 24 are mated using an interbody piece 30. As can be seen in Figs. 3a, 3b, the first forked end 22 includes a threaded shaft 32 which projects rearwardly

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from the saddle 26. The threaded shaft 32 fits into the threaded bore 34 (Fig. 4a) of the interbody piece 30.

5 The second forked end 24 (Figs. 5a, 5b) includes a smooth cylindrical shaft 36 which can fit into the smooth bore 38 of the interbody piece 30.

Fig. 1 shows the implant 20 in a fully extended position, while Fig. 2 shows the implant in an unextended position. In the unextended position, it can be seen that the threaded shaft 32 of the first forked end 22 fits inside the hollow cylindrical shaft 36 of the second forked end 24.

10 For purposes of implantation between adjacent first and second spinous processes of the spinal column, the implant 20 is configured as shown in Fig. 2. The first and second spinous processes are exposed using appropriate surgical techniques and thereafter, the implant 20 is positioned so that saddle 26 engages the first spinous process, and
15 saddle 28 engages the second spinous process. At this point, the interbody piece 30 can be rotated by placing an appropriate tool or pin into the cross holes 40 and upon rotation, the saddle 26 is moved relative to the saddle 28. Such rotation spreads apart or distracts the spinous processes with the resultant and beneficial effect of enlarging
20 the volume of the spinal canal in order to alleviate any restrictions on blood vessels and nerves.

It is noted that this implant as well as the several other implants described herein act as an extension stop. That means that as the back is bent backwardly and thereby placed in extension the spacing between
25 adjacent spinous processes cannot be reduced to a distance less than the distance between the lowest point of saddle 26 and the lowest point of saddle 28. This implant, however, does not inhibit or in any way limit the flexion of the spinal column, wherein the spinal column is bent forward.

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Preferably, such a device provides for distraction in the range of about 5 mm to about 15 mm. However, devices which can distract up to and above 22 mm may be used depending on the characteristics of the individual patient.

5 With all the ligaments (such as the supraspinous ligament) and tissues associated with the spinous processes left intact, the implant 20 can be implanted essentially floating in position in order to gain the benefits of the aforementioned extension stop and flexion non-inhibitor. If desired, one of the saddles 26 can be laterally pinned with pin 29 to
10 one of the spinous processes and the other saddle can be loosely associated with the other spinous processes by using a tether 31 which either pierces or surrounds the other spinous process and then is attached to the saddle in order to position the saddle relative to the spinous process. Alternatively, both saddles can be loosely tethered to
15 the adjacent spinous process in order to allow the saddles to move relative to the spinous processes.

The shape of the saddles, being concave, gives the advantage of distributing the forces between the saddle and the respective spinous process. This ensures that the bone is not resorbed due to the
20 placement of the implant 20 and that the structural integrity of the bone is maintained.

The implant 20 in this embodiment can be made of a number of materials, including but not limited to, stainless steel, titanium, ceramics, plastics, elastics, composite materials or any combination of the above.
25 In addition, the modulus of elasticity of the implant can be matched to that of bone, so that the implant 20 is not too rigid. The flexibility of the implant can further be enhanced by providing additional apertures or perforations throughout the implant in addition to the holes 40 which

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also have the above stated purpose of allowing the interbody piece 30 to be rotated in order to expand the distance between the saddle 26, 28.

In the present embodiment, it is understood that the spinous processes can be accessed and distracted initially using appropriate instrumentation, and that the implant 20 can be inserted and adjusted
5 in order to maintain and achieve the desired distraction. Alternatively, the spinous process can be accessed and the implant 20 appropriately positioned. Once positioned, the length of the implant can be adjusted in order to distract the spinous processes or extend the distraction of
10 already distracted spinous processes. Thus, the implant can be used to create a distraction or to maintain a distraction which has already been created.

The placement of implants such as implant 20 relative to the spinous process will be discussed hereinbelow with other embodiments.
15 However, it is to be noted that ideally, the implant 20 would be placed close to the instantaneous axis of rotation of the spinal column so that the forces placed on the implant 20 and the forces that the implant 20 places on the spinal column are minimized.

Further, it is noted that during the actual process of installing or
20 implanting the implant 20, that the method uses the approach of extending the length of the implant 20 a first amount and then allowing the spine to creep or adjust to this distraction. Thereafter, implant 20 would be lengthened another amount, followed by a period where the spine is allowed to creep or adjust to this new level of distraction. This
25 process could be repeated until the desired amount of distraction has been accomplished. This same method can be used with insertion tools prior to the installation of an implant. The tools can be used to obtain the desired distraction using a series of spinal distraction and spine creep periods before an implant is installed.

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Embodiment of Figs. 6, 7, 8, 9 and 10

The embodiment of the invention shown in the above Figs. 6, 7, 8, 9 and 10 includes distraction or spreader tool 50 which has first and second arms 52, 54. Arms 52, 54 are pivotal about pivot point 56 and
5 releaseable from pivot point 56 in order to effect the implantation of implant 58. As can be seen in Fig. 6, in cross-section, the arms 52, 54 are somewhat concave in order to cradle and securely hold the first spinous process 60 relative to arm 52 and the second spinous process 62 relative to arm 54. The distraction tool 50 can be inserted through a
10 small incision in the back of the patient in order to address the space between the first spinous process 60 and the second spinous process 62. Once the tool 50 is appropriately positioned, the arms 52, 54 can be spread apart in order to distract the spinous processes. After this has occurred, an implant 58 as shown in Figs. 8 and 9, or of a design shown
15 in other of the embodiments of this invention, can be urged between the arms 52, 54 and into position between the spinous processes. After this occurs, the arms 52, 54 can be withdrawn from the spinous processes leaving the implant 58 in place. The implant 58 is urged into place using a tool 64 which can be secured to the implant 58 through a threaded
20 bore 66 in the back of the implant. As can be seen in Fig. 10, the implant 58 includes saddles 68 and 70 which cradle the upper and lower spinous processes 60, 62 in much the same manner as the above first embodiment and also in much the same manner as the individual arms of the tool 50. The saddles as described above tend to distribute the
25 load between the implant and the spinous processes and also assure that the spinous process is stably seated at the lowest point of the respective saddles.

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Embodiment of Figs. 11, 12 and 13

Another embodiment of the apparatus and method of the invention is shown in Figs. 11, 12 and 13. In this embodiment, the spreader or distraction tool 80 includes first and second arms 82, 84 which are permanently pivoted at pivot point 86. The arms include L-shaped ends 88, 90. Through a small incision, the L-shaped ends 88, 90 can be inserted between the first and second spinous processes 92, 94. Once positioned, the arms 82, 84 can be spread apart in order to distract the spinous processes. The implant 96 can then be urged between the spinous processes in order to maintain the distraction. It is noted that implant 96 includes wedged surfaces or ramps 98, 100. As the implant 96 is being urged between the spinous processes, the ramps further cause the spinous processes to be distracted. Once the implant 96 is fully implanted, the full distraction is maintained by the planar surfaces 99, 101 located rearwardly of the ramps. It is to be understood that the cross-section of the implant 96 can be similar to that shown for implant 58 or similar to other implants in order to gain the advantages of load distribution and stability.

Embodiments of Figs. 14, 15, 16, 16a, and 17

In Figs. 14 and 15, yet another embodiment of the invention is depicted. In this embodiment, the implant 110 includes first and second conically shaped members 112, 114. Member 112 includes a male snap connector 116 and member 114 includes a female snap connector 118. With male snap connector 116 urged into female snap connector 118, the first member 112 is locked to the second member 114. In this embodiment, a distraction or spreader tool 80 could be used. Once the spinous process has been spread apart, an implantation tool 120 can be used to position and snap together the implant 110. The first member

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112 of implant 110 is mounted on one arm and second member 114 is mounted on the other arm of tool 120. The member 112, 114 are placed on opposite sides of the space between adjacent spinous processes. The members 112, 114 are urged together so that the
5 implant 110 is locked in place between the spinous processes as shown in Fig. 15. It is to be noted that the implant 110 can also be made more self-distracting by causing the cylindrical surface 122 to be more conical, much as surface 124 is conical, in order to hold implant 110 in place relative to the spinous processes and also to create additional
10 distraction.

An alternative embodiment of the implant can be seen in Figs. 16 and 17. This implant 130 includes first and second members 132, 134. In this particular embodiment, the implants are held together using a screw (not shown) which is inserted through countersunk bore 136 and
15 engages a threaded bore 138 of the second member 134. Surfaces 139 are flattened (Fig. 17) in order to carry and spread the load applied thereto by the spinous processes.

The embodiment of implant 130 is not circular in overall outside appearance, as is the embodiment 110 of Figs. 14 and 15. In particular,
20 with respect to the embodiment of implant 130 of Figs. 16 and 17, this embodiment is truncated so that the lateral side 140, 142 are flattened with the upper and lower sides 144, 146 being elongated in order to capture and create a saddle for the upper and lower spinous processes. The upper and lower sides, 144, 146 are rounded to provide a more
25 anatomical implant which is compatible with the spinous processes.

If it is desired, and in order to assure that the first member 132 and the second member 134 are aligned, key 148 and keyway 150 are designed to mate in a particular manner. Key 148 includes at least one flattened surface, such as flattened surface 152, which mates to an

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appropriately flattened surface 154 of the keyway 150. In this manner, the first member is appropriately mated to the second member in order to form appropriate upper and lower saddles holding the implant 130 relative to the upper and lower spinous processes.

5 Fig. 16a depicts second member 134 in combination with a rounded nose lead-in plug 135. Lead-in plug 135 includes a bore 137 which can fit snugly over key 148. In this configuration, the lead-in plug 135 can be used to assist in the placement of the second member 134 between spinous processes. Once the second member 134 is
10 appropriately positioned, the lead-in plug 135 can be removed. It is to be understood that the lead-in plug 135 can have other shapes such as pyramids and cones to assist in urging apart the spinous processes and soft tissues in order to position the second member 134.

15 **Embodiment of Figs. 18, 19 and 20**

 The implant 330 as shown in Fig. 18 is comprised of first and second mating wedges 332 and 334. In order to implant these wedges 332, 334, the spinous processes are accessed from both sides and then a tool is used to push the wedges towards each other. As the wedges
20 are urged towards each other, the wedges move relative to each other so that the combined dimension of the implant 330 located between the upper and lower spinous processes 336, 338 (Fig. 20), increases, thereby distracting the spinous processes. It is noted that the wedges 332, 334 include saddle 340, 342, which receiving the spinous
25 processes 336, 338. These saddles have the advantages as described hereinabove.

 The first or second wedges 332, 334 have a mating arrangement which includes a channel 344 and a projection of 346 which can be urged into the channel in order to lock the wedges 332, 334 together.

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The channel 334 is undercut in order to keep the projection from separating therefrom . Further, as in other devices described herein, a detent can be located in one of the channel and the projection, with a complimentary recess in the other of the channel and the projection.
5 Once these two snap together, the wedges are prevented from sliding relative to the other in the channel 344.

While the above embodiment was described with respect to wedges, the wedges could also have been designed substantially as cones with all the same features and advantages.

10

Embodiments of Figs. 21 and 22

The implant 370 is comprised of first and second distraction cone 372, 374. These cones are made of a flexible material. The cones are positioned on either side of the spinous processes 376, 378 as shown
15 in Fig. 21. Using appropriate tool as shown hereinabove, the distraction cones 372, 374 are urged together. As they are urged together, the cones distract the spinous processes as shown in Fig. 22. Once this has occurred, an appropriate screw or other type of fastening mechanism 380 can be used to maintain the position of the distraction
20 cones 372, 374. The advantage of this arrangement is that the implant 370 is self-distracting and also that the implant, being flexible, molds about the spinous processes as shown in Fig. 22.

Embodiments of Fig. 23, 24 and 25

25 In Figs. 23 and 24, another embodiment of the implant 170 is depicted. This implant is guided in place using an L-shaped guide 172 which can have a concave cross-section such as the cross-section 52 of retraction tool 50 in Fig. 6 in order to cradle and guide the implant 170 in position. Preferably a small incision would be made into the back of

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the patient and the L-shaped guide tool 172 inserted between the adjacent spinous processes. The implant 170 would be mounted on the end of insertion tool 174 and urged into position between the spinous processes. The act of urging the implant into position could cause the spinous processes to be further distracted if that is required. Prior to the
5 insertion of the L-shaped guide tool 172, a distraction tool such as shown in Fig. 13 could be used to initially distract the spinous processes.

Implant 170 can be made of a deformable material so that it can
10 be urged into place and so that it can somewhat conform to the shape of the upper and lower spinous processes. This deformable material would be preferably an elastic material. The advantage of such a material would be that the load forces between the implant and the spinous processes would be distributed over a much broader surface
15 area. Further, the implant would mold itself to an irregular spinous process shape in order to locate the implant relative to spinous processes.

With respect to Fig. 25, this implant 176 can be inserted over a guide wire, guide tool or stylet 178. Initially, the guide wire 178 is
20 positioned through a small incision to the back of the patient to a position between the adjacent spinous processes. After this has occurred, the implant is threaded over the guide wire 178 and urged into position between the spinous processes. This urging can further distract the spinous processes if further distraction is required. Once the implant
25 is in place, the guide tool 178 is removed and the incision closed. The insertion tools of Figs. 23 and 24 can also be used if desired.

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Embodiment of Figs. 26, 27 and 28

The embodiment shown in Figs. 26, 27 and 28 uses an implant similar to that depicted in Figs. 8 and 9 with different insertion tools. As can be seen in Fig. 26, an L-shaped distraction tool 190 is similar to L-shaped distraction tool 80 (Fig. 12), is used to distract the first and second spinous processes 192, 194. After this has occurred, an insertion tool 196 is placed between the spinous processes 192, 194. Insertion tool 196 includes a handle 198 to which is mounted a square-shaped ring 200.

The distraction tool 190 can be inserted through a small incision in the back in order to spread apart the spinous processes. Through the same incision which has been slightly enlarged laterally, an upper end 202 of ring 200 can be initially inserted followed by the remainder of the ring 200. Once the ring is inserted, the ring can be rotated slightly by moving handle 198 downwardly in order to further wedge the spinous processes apart. Once this has been accomplished, an implant such as implant 204 can be inserted through the ring and properly positioned using implant handle 206. Thereafter, the implant handle 206 and the insertion tool 196 can be removed.

Embodiments of Figs. 29, 30, 31, 32 and 33

As can be seen in Figs. 29 and 30, the implants 210, 212, can have different shapes when viewed from the side. These implants are similar to the above-referenced implants 58 (Fig. 8) and 204 (Fig. 28). These implants have cross-sections similar to that shown in Fig. 10 which includes saddles in order to receive and hold the adjacent spinous processes.

As can be seen in Figs. 31, 32 and 33, these implants can be placed in different positions with respect to the spinous process 214.

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Preferably as shown in Fig. 33, the implant 210 is placed closest to the lamina 216. Being so positioned, the implant 210 is close to the instantaneous axis of rotation 218 of the spinal column, and the implant would experience least forces caused by movement of the spine. Thus, 5 theoretically, this is the optimal location for the implant.

As can be seen in Figs. 31 and 32, the implant can be placed midway along the spinous process (Fig. 32) and towards the posterior aspect of the spinous process (Fig. 31). As positioned shown in Fig. 31, the greatest force would be placed on the implant 210 due to a 10 combination of compression and extension of the spinal column.

Embodiment of Figs. 34 and 35

Another embodiment of the invention is shown in Figs. 34 and 35. In these figures, implant 220 is comprised of a plurality of individual 15 leaves 222 which are substantially V-shaped. The leaves include interlocking indentations or detents 224. That is, each leaf includes an indentation with a corresponding protrusion such that a protrusion of one leaf mates with an indentation of an adjacent leaf. Also associated with this embodiment is an insertion tool 226 which has a blunt end 228 20 which conforms to the shape of an individual leaf 222. For insertion of this implant into the space between the spinous processes as shown in Fig. 29, the insertion tool 226 first insert a single leaf 220. After that has occurred, the insertion tool then inserts a second leaf with the protrusion 224 of the second leaf snapping into corresponding indentation made 25 by the protrusion 224 of the first leaf. This process would reoccur with third and subsequent leaves until the appropriate spacing between the spinous processes was built up. As can be seen in Fig. 29, the lateral edges 229 of the individual leaves 222 are slightly curved upwardly in

- 25 -

order to form a saddle for receiving the upper and lower spinous processes.

Embodiments of Figs. 36, 37 and 38

5 The embodiments of Figs. 36, 37 and 38 which include implants 230, 232, and 234 respectively, are designed in such a manner so the implant locks itself into position once it is properly positioned between the spinous processes. Implant 220 is essentially a series of truncated cones and includes a plurality of ever expanding steps 236. These
10 steps are formed by the conical bodies starting with the nose body 238 followed there behind by conical body 240. Essentially, the implant 234 looks like a fir tree placed on its side.

 The implant 230 is inserted laterally throughout the opening between upper and lower spinous processes. The first body 238 causes
15 the initial distraction. Each successive conical body distracts the spinous processes a further incremental amount. When the desired distraction has been reached, the spinous processes are locked into position by steps 236. At this point, if desired, the initial nose body 238 of the implant and other bodies 240 can be broken, snapped or sawed
20 off if desired in order to minimize the size of the implant 230. In order for a portion of the implant 230 to be broken or snapped off, the intersection between bodies such as body 238 and 240, which is intersection line 242, would be somewhat weakened with the appropriate removal of material. It is noted that only the intersection lines of the
25 initial conical bodies need to be so weakened. Thus, intersection line 244 between the bodies which remain between the spinous processes would not need to be weaker, as there would be no intention that the implant would be broken off at this point.

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Fig. 37 shows implant 232 positioned between upper and lower spinous processes. This implant is wedge-shaped or triangular shaped in cross-sectioned and includes bore pluralities 245 and 246. Through these bores can be placed locking pins 248 and 250. The triangular or
5 wedged-shaped implant can be urged laterally between and thus distract the upper and lower spinous processes. Once the appropriate distraction is reached, pins 248, 250 can be inserted through the appropriate bores of the bore pluralities 245 and 246 in order to lock the spinous processes in a V-shaped valley formed by pins 248, 250 on the
10 one hand and the ramped surface 233, 235 on the other hand.

Turning to Fig. 38, the implant 234 has a triangular-shaped or wedge-shaped body similar to that shown in Fig. 32. In this embodiment, tab 252, 254 are pivotally mounted to the triangular shaped body 234. Once the implant 234 is appropriately positioned in
15 order to distract the spinous processes to the desired amount, the tabs 252, 254 rotate into position in order to hold the implant 234 in the appropriate position.

Embodiment of Figs. 39 and 40

20 In the embodiment of Figs. 39 and 40, cannula 258 is inserted through a small incision to a position between upper and lower spinous processes. Once the cannula is properly inserted, an implant 260 is pushed through the cannula 258 using an insertion tool 262. The implant 260 includes a plurality of ribs or indentation 264 that assist in
25 positioning the implant 260 relative to the upper and lower spinal processes. Once the implant 260 is in position, the cannula 258 is withdrawn so that the implant 260 comes in contact with and wedges between the spinous processes. The cannula 258 is somewhat conical in shape with the nose end 266 being somewhat smaller than the distal

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end 268 in order to effect the insertion of the cannula into the space between the spinous processes.

Further, a plurality of cannula can be used instead of one, with each cannula being slightly bigger than one before. In the method of the invention, the first smaller cannula would be inserted followed by successively larger cannula being placed over the previous smaller cannula. The smaller cannula would then be withdrawn from the center of the larger cannula. Once the largest cannula is in place, and the opening of the skin accordingly expanded, the implant, which is accommodated by only the larger cannula, is inserted through the larger cannula and into position.

Embodiments of Figs. 41, 42 and 43

The precurved implant 270 in Figs. 41 and 42, and precurved implant 272 in Fig. 43 have common introduction techniques which includes a guide wire, guide tool, or stylet 274. For both embodiments, the guide wire 274 is appropriately positioned through the skin of the patient and into the space between the spinous processes. After this is accomplished, the implant is directed over the guide wire and into position between the spinous processes. The precurved nature of the implant assist in (1) positioning the implant through a first small incision in the patient's skin on one side of the space between two spinous processes and (2) guiding the implant toward a second small incision in the patient's skin on the other side of the space between the two spinous processes. With respect to the implant 270, the implant includes a conical introduction nose 276 and a distal portion 278. As the nose 276 is inserted between the spinous processes, this causes distraction of the spinous processes. Break lines 280, 282 are established at opposite sides of the implant 270. Once the implant is

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properly positioned over the guide wire between the spinous processes, the nose portion 276 and the distal portion 278 can be broken off along the break lines, through the above two incisions, in order to leave the implant 270 in position.

5 Although only two break lines 280, 282 are depicted, multiple break lines can be provided on implant 270 so that the implant can continue to be fed over the guide wire 278 until the appropriate width of the implant 270 creates the desired amount of distraction. As described hereinabove, the break lines can be created by perforating or otherwise
10 weakening the implant 270 so that the appropriate portions can be snapped or sawed off.

 With respect to the precurved implant 272, this implant is similar in design to the implant 230 shown in Fig. 36. This implant 272 in Fig. 47, however, is precurved and inserted over a guide wire 274 to a
15 position between the spinous processes. As with implant 230 in Fig. 43, once the appropriate level of this distraction has been reached and if desired, sections of the implant 272 can be broken, snapped or sawed off as described hereinabove in order to leave a portion of the implant wedged between the upper and lower spinous processes.

20

Embodiment of Fig. 44

 A further embodiment of the invention is shown in Fig. 44. This embodiment includes a combination insertion tool and implant 290. The insertion tool and implant 290 is in the shape of a ring which is hinged
25 at point 292. The ring is formed by a first elongated and conically shaped member 294 and a second elongated and conically shaped member 296. Members 294 and 296 terminate in points and through the use of hinge 292 are aligned and meet. Through similar incisions on both sides of the spinous processes, first member and second member

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are inserted through the skins of the patient and are mated together between the spinous processes. After this has occurred, the implant 290 is rotated, for example clockwise, so that increasingly widening portions of the first member 292 are used to distract the first and second spinous processes. When the appropriate level of distraction has occurred, the remainder of the ring before and after the section which is located between the spinous processes can be broken off as taught hereinabove in order to maintain the desired distraction. Alternatively, with a small enough ring, the entire ring can be left in place with the spinous processes distracted.

Embodiment of Fig. 45

In Fig. 45, the implant 300 is comprised of a plurality of rods or stylets 302 which are inserted between the upper and lower spinous processes. The rods are designed much as described hereinabove so that they may be broken, snapped or cut off. Once these are inserted and the appropriate distraction has been reached, the stylets are broken off and a segment of each stylet remains in order to maintain distraction of the spinous process.

Embodiment of Figs. 46 and 47

Implant 310 of Figs. 46 and 47 is comprised of a shape memory material which coils upon being released. The material is straightened out in a delivery tool 312. The delivery tool is in position between upper and lower spinous processes 314, 316. The material is then pushed through the delivery tool. As it is released from the delivery end 318 of the delivery tool, the material coils, distracting the spinous processes to the desired amount. Once this distraction has been achieved, the material is cut and the delivery tool removed.

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Embodiments of Figs. 48, 49, 50 and 51

As can be seen in Fig. 48, the implant 320 is delivered between upper and lower spinous processes 322 and 324, by delivery tool 326. Once the implant 320 is in place between the spinous processes, the
5 delivery tool is given a 90° twist so that the implant goes from the orientation as shown in Fig. 49, with longest dimension substantially perpendicular to the spinous processes, to the orientation shown in Fig. 50 where the longest dimension is in line with and parallel to the spinous processes. This rotation causes the desired distraction between the
10 spinous processes. Implant 320 includes opposed recesses 321 and 323 located at the ends thereof. Rotation of the implant 320 causes the spinous processes to become lodged in these recesses.

Alternatively, the insertion tool 326 can be used to insert multiple implants 320, 321 into the space between the spinous processes 322,
15 324 (Fig. 51). Multiple implants 320, 321 can be inserted until the appropriate amount of distraction is built up. It is to be understood in this situation that one implant would lock to another implant by use of, for example, a channel arrangement wherein a projection from one of the implants would be received into and locked into a channel of the
20 other implant. Such a channel arrangement is depicted with respect to the other embodiment.

Embodiment of Figs. 52, 53, 54, 55a and 55b

The embodiment of Figs. 52 through 55b is comprised of a fluid-
25 filled dynamic distraction implant 350. This implant includes a membrane 352 which is placed over pre-bent insertion rod 354 and then inserted through an incision on one side of the spinous process 356. The bent insertion rod, with the implant 350 thereover, is guided between appropriate spinous processes. After this occurs, the insertion

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rod 354 is removed leaving the flexible implant in place. The implant 350 is then connected to a source of fluid (gas, liquid, gel and the like) and the fluid is forced into the implant causing it to expand as shown in Fig. 54, distracting the spinal processes to the desired amount. Once
5 the desired amount of distraction has occurred, the implant 350 is closed off as is shown in Fig. 55a. The implant 350 being flexible, can mold to the spinous processes which may be of irregular shape, thus assuring positioning. Further, implant 350 acts as a shock absorber, damping forces and stresses between the implant and the spinous
10 processes.

A variety of materials can be used to make the implant and the fluid which is forced into the implant. By way of example only, viscoelastic substances such as methylcellulose, or hyaluronic acid can be used to fill the implant. Further, materials which are initially a fluid,
15 but later solidify, can be inserted in order to cause the necessary distraction. As the materials solidify, they mold into a custom shape about the spinous processes and accordingly are held in position at least with respect to one of two adjacent spinous processes. Thus, it can be appreciated that using this embodiment and appropriate insertion
20 tools the implant can be formed about one spinous process in such a manner that the implant stays positioned with respect to that spinous process (Fig. 55b). With such an embodiment, a single implant can be used as an extension stop for spinous process located on either side, without restricting flexion of the spinal column.

25 It is to be understood that many of the other implants disclosed herein can be modified so that they receive a fluid in order to establish and maintain a desired distraction much in the manner as implant 350 receives a fluid.

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Embodiment of Figs. 56, 57 and 58

The implant 360 as shown in Fig. 56 is comprised of a shape memory material such as a plastic or a metal. A curved introductory tool 362 is positioned between the appropriate spinous processes as described hereinabove. Once this has occurred, bore 364 of the implant is received over the tool. This act can cause the implant to straighten out. The implant is then urged into position and thereby distracts the spinous processes. When this has occurred, the insertion tool 362 is removed, allowing the implant to assume its pre-straightened configuration and is thereby secured about one of the spinous processes. Such an arrangement allows for an implant that is an extension stop and does not inhibit flexion of the spinous column. Alternatively, the implant can be temperature sensitive. That is to say that the implant would be more straightened initially, but become more curved when it was warmed by the temperature of the patient's body.

Embodiments of Figs. 59 and 60

In this embodiment, the implant 380 is comprised of a plurality of interlocking leaves 382. Initially, a first leaf is positioned between opposed spinous processes 384, 386. Then subsequently, leaves 382 are interposed between the spinous processes until the desired distraction has been built up. The leaves are somewhat spring-like in order to absorb the shock and can somewhat conform to the spinous processes.

Embodiment of Fig. 61

The implant 390 of Fig. 61 includes the placement of shields 392, 394 over adjacent spinous processes 396, 398. The shields are used to prevent damage to the spinous processes. These shields include apertures which receives a self-tapping screw 400, 402. In practice, the

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shields are affixed to the spinous processes and the spinous processes are distracted in the appropriate amount. Once this has occurred, a rod 404 is used to hold the distracted position by being screwed into each of the spinous processes through the aperture in the shields using the screws as depicted in Fig. 61.

Embodiment of Figs. 62 and 63

Implant 410 of Figs. 62, 63 is comprised of first and second members 412, 414 which can be mated together using an appropriate screw and threaded bore arrangement to form the implant 410. Main member 412 and mating member 414 form implant 410. Accordingly, the implant 410 would have a plurality of members 414 for use with a standardized first member 412. Figs. 62 and 64 show different types of mating members 414. In Fig. 62, the mating member 414 includes projections 416 and 418 which act like shims. These projections are used to project into the space of saddles 420, 422 of the first member 412. These projections 416, 418 can be of varying lengths in order to accommodate different sizes of spinous processes. A groove 424 is placed between the projections 416, 418 and mates with an extension 426 of the first member 412.

As shown in Fig. 63, the projections of the embodiment shown in Fig. 62 are removed and recesses 428, 430 are substituted therefor. These recesses expand the area of the saddles 420, 422 in order to accommodate larger spinous processes.

Embodiment of Figs. 64, 65 and 66

The embodiments of Figs. 64, 65 and 66 are similar in design and concept to the embodiment of Figs. 62 and 63. In Fig. 64, the implant 500 includes the first and second members 502, 504. These members

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can be secured together with appropriate screws or other fastening means as taught in other embodiments. Implant 500 includes first and second saddles 506, 508 which are formed between the ends of first and second members 502, 504. These saddles 506, 508 are used to receive and cradle the adjacent spinous processes. As can be seen in Fig. 64, each saddle 506, 508 is defined by a single projection or leg 510, 512, which extends from the appropriate first and second members 502, 504. Unlike the embodiment found in Figs. 62 and 63, each of the saddles is defined by only a single leg as the ligaments and other tissues associated with the spinous processes can be used to ensure that the implant is held in an appropriate position. With the configuration of Fig. 64, it is easier to position the implant relative to the spinous processes as each saddle is defined by only a single leg and thus the first and second members can be more easily worked into position between the various tissues.

In the embodiment of Fig. 65, the implant 520 is comprised of a single piece having saddles 522 and 524. The saddles are defined by a single leg 526, 528 respectively. In order for this implant 520 to be positioned between the spinous processes, an incision is made between lateral sides of adjacent spinous processes. The single leg 526 is directed through the incision to a position adjacent to an opposite lateral side of the spinous process with the spinous process cradled in the saddle 522. The spinous processes are then urged apart until saddle 524 can be pivoted into position into engagement with the other spinous process in order to maintain the distraction between the two adjacent spinous processes.

The embodiment of Fig. 66 is similar to that of Fig. 65 with an implant 530 and first and second saddles 532 and 534. Associated with each saddle is a tether 536, 538 respectively. The tethers are made of

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flexible materials known in the trade and industry and are positioned through bores in the implant 530. Once appropriately positioned, the tethers can be tied off. It is to be understood that the tethers are not meant to be used to immobilize one spinous process relative to the other, but are used to guide motion of the spinous processes relative to each other so that the implant 530 can be used as an extension stop and a flexion non-inhibitor. In other words, the saddles 532, 534 are used to stop spinal column backward bending and extension. However, the tethers do not inhibit forward bending and spinal column flexion.

10

Embodiments of Figs. 67, 68

The implant 550 is Z-shaped and includes a central body 552 and first and second arms 554, 556, extending in opposite directions therefrom. The central body 552 of the implant 550 includes first and second saddles 558 and 560. The first and second saddles 558 and 560 would receive upper and lower spinous processes 562, 568. The arms 554, 556 are accordingly located adjacent the distal end 566 (Fig. 68) of the central body 552. The first and second arms 554, 556, act to inhibit forward movement, migration or slippage of the implant 550 toward the spinal canal and keep the implant in place relative to the first and second spinal processes. This prevents the implant from pressing down on the ligamentum flavum and the dura. In a preferred embodiment, the central body would have a height of about 10mm with each of the arms 554, 556 have a height of also about 10mm. Depending on the patient, the height of the body could vary from about less than 10 mm to about greater than 24mm. As can be seen in Figs. 67 and 68, the first and second arms 554, 556 are additionally contoured in order to accept the upper and lower spinous processes 556, 558. In particular, the arms 554, 556 as can be seen with respect

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to arm 554 have a slightly outwardly bowed portion 568 (Fig. 68) with a distal end 570 which is slightly inwardly bowed. This configuration allows the arm to fit about the spinous process with the distal end 570 somewhat urged against the spinous process in order to guide the motion of the spinous process relative to the implant. These arms 554, 556 could if desired to be made more flexible than the central body 552 by making arms 554, 556 thin and/or with perforations, and/or other material different than that of the central body 550. As with the last embodiment, this embodiment can be urged into position between adjacent spinous processes by directing an arm into a lateral incision so that the central body 552 can be finally positioned between spinous processes.

Embodiment of Figs. 69, 70, 71 and 71a

Figs. 69, 70 and 71 are perspective front, end, and side views of implant 580 of the invention. This implant includes a central body 582 which has first and second saddles 584, 586 for receiving adjacent spinous processes. Additionally, the implant 580 includes first and second arms 588 and 590. The arms, as with the past embodiment, prevent forward migration or slippage of the implant toward the spinal canal. First arm 588 projects outwardly from the first saddle 584 and second arm 590 projects outwardly from the second saddle 586. In a preferred embodiment, the first arm 588 is located adjacent to the distal end 600 of the central body 582 and proceeds only partly along the length of the central body 582. The first arm 588 is substantially perpendicular to the central body as shown in Fig. 70. Further, the first arm 588, as well as the second arm 590, is anatomically rounded.

The second arm 590, projecting from second saddle 586, is located somewhat rearward of the distal end 600, and extends partially

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along the length of the central body 582. The second arm 590 projects at a compound angle from the central body 582. As can be seen in Figs. 70 and 71, the second arm 590 is shown to be at about an angle of 45° from the saddle 586 (Fig. 70). Additionally, the second arm 590 is at an angle of about 45° relative to the length of the central body 580 as shown in Fig. 71. It is to be understood that other compound angles are within the spirit and scope of the invention as claimed.

In a preferred embodiment, the first and second arms 588, 590 have a length which is about the same as the width of the central body 582. Preferably, the length of each arm is about 10mm and the width of the central body is about 10mm. However, the bodies with the widths of 24mm and greater are within the spirit and scope of the invention, along with first and second arms ranging from about 10mm to greater than about 24mm. Further, it is contemplated that the embodiment could include a central body having a width of about or greater than 24mm with arms being at about 10mm.

It is to be understood that the embodiment of Figs. 69, 70 and 71 as well as the embodiment of Figs. 67 and 68 are designed to preferably be positioned between the L4-L5 and the L5-S1 vertebral pairs. The embodiment of Figs. 69, 70, 71 is particularly designed for the L5-S1 position with the arms being designed to conform to the sloping surfaces found therebetween. The first and second arms are thus contoured so that they lie flat against the lamina of the vertebra which has a slight angle.

The embodiment of Fig. 69, 70, and 71 as with the embodiment of Figs. 67 and 68 is Z-shaped in configuration so that it may be inserted from one lateral side to a position between adjacent spinous processes. A first arm, followed by the central body, is guided through the space between the spinous processes. Such an arrangement only requires

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that an incision on one side of the spinous process be made in order to successfully implant the device between the two spinous processes.

5 The implant 610 of Fig. 71a is similar to that immediately above with the first arm 612 located on the same side of the implant as the second arm 614. The first and second saddle 616, 618 are slightly modified in that distal portion 620, 622 are somewhat flattened from the normal saddle shape in order to allow the implant to be positioned between the spinous processes from one side. Once in position, the ligaments and tissues associated with the spinous processes would hold
10 the implant into position. Tethers also could be used if desired.

Embodiment of Figs. 72, 73

15 Implant 630 is also designed so that it can be inserted from one side of adjacent spinous processes. This insert 630 includes a central body 632 with the first and second arms 634, 636 extending on either side thereof. As can be seen in Fig. 72, a plunger 638 is positioned to extend from an end of the central body 632. As shown in Fig. 72, the plunger 638 is fully extended and as shown in Fig. 73, the plunger 638 is received within the central body 632 of the implant 630. With the
20 plunger received into the implant 632, the third and fourth arms or hooks 640, 642 can extend outwardly from the central body 632. The third and fourth arms or hooks 640, 642 can be comprised of a variety of materials, such as for example, shape memory metal materials or materials which have a springy quality.

25 For purposes of positioning the implant 630 between adjacent spinous processes, the plunger 638 is pulled outwardly as shown in Fig. 72. The central body 632 is then positioned between adjacent spinous processes and the plunger 638 is allowed to move to the position of Fig. 73 so that the third and fourth arms 640, 642 can project

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outwardly from the central body 632 in order to hold the implant 630 in position between the spinous processes.

Plunger 638 can be spring biased to the position as shown in Fig. 73 or can include detents or other mechanisms which lock it into that position. Further, the third and fourth arms themselves, as deployed, can keep the plunger in the position as shown in Fig. 73.

Embodiments of Figs. 74, 75, 76, 77, and 78

Other embodiments of the invention are shown in Figs. 74 through 78. Figs. 74, 75 and 76 disclose implant 700. Implant 700 is particularly suited for implantation between the L4-L5 and L5-S1 vertebra. As can be seen in Fig. 74, the implant 700 includes a central body 702 which has a bore 704 provided therein. Bore 704 is used in order to adjust the modulus of elasticity of the implant so that it is preferably approximately two times the anatomical load placed on the vertebra in extension. In other words, the implant 700 is approximately two times stiffer than the normal load placed on the implant. Such an arrangement is made in order to ensure that the implant is somewhat flexible in order to reduce potential resorption of the bone adjacent to the implant. Other modulus values can be used and be within the spirit of the invention.

Implant 700 includes first and second saddle 706, 708 which are used to receive and spread the load from the upper and lower spinous processes. The saddle 706 is defined by first and second arms 710 and 712. The second saddle 708 is defined by third and fourth arms 714 and 716. As can be seen in Fig. 74, the first arm 710, in a preferred embodiment, is approximately two times the length of the body 702 with the second arm being approximately less than a quarter length of the body. Third arm 714 is approximately one times the length of the body

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702 with the fourth arm 716 being, in this preferred embodiment, approximately one and a half times the length of the body 702. The arms are designed in such a way that the implant (1) can be easily and conveniently inserted between the adjacent spinous processes, (2) will
5 not migrate forwardly toward the spinal canal, and (3) will hold its position through flexion and extension as well as lateral bending of the spinal column.

First arm 710 is in addition designed to accommodate the shape of the vertebra. As can be seen in Fig. 74, the first arm 710 becomes
10 narrower as it extends away from the body 702. The first arm 710 includes a sloping portion 718 followed by a small recess 720 ending in a rounded portion 722 adjacent to the end 724. This design is provided to accommodate the anatomical form of for example the L4 vertebra. It is to be understood that these vertebra have a number of surfaces at
15 roughly 30° angles and that the sloping surfaces of this embodiment and the embodiments shown in Figs. 77 and 78 are designed to accommodate these surfaces. These embodiments can be further modified in order to accommodate other angles and shapes.

The second arm 712 is small so that it is easy to insert between
20 the spinous processes, yet still define the saddle 706. The fourth arm 716 is larger than the third arm 714, both of which are smaller than the first arm 710. The third and fourth arms are designed so that they define the saddle 706, guide the spinous processes relative to the implant 700 during movement of the spinal column, and yet are of a size
25 which makes the implant easy to position between the spinous processes.

The procedure, by way of example only, for implanting the implant 700 can be to make an incision laterally between two spinous processes and then initially insert first arm 710 between the spinous processes.

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The implant and/or appropriate tools would be used to distract the spinous processes allowing the third leg 714 and the central body 702 to fit through the space between the spinous processes. The third leg 714 would then come to rest adjacent the lower spinous processes on the opposite side with the spinous processes resting in the first and second saddle 706, 708. The longer fourth leg 716 would then assist in the positioning of the implant 700.

Fig. 77 includes an implant 740 which is similar to implant 700 and thus have similar numbering. The saddle 706, 708 of implant 740 have been cantered or sloped in order to accommodate the bone structure between, by way of example, the L4-L5 and the L5-S1 vertebra. As indicated above, the vertebra in this area have a number of sloping surfaces in the range of about 30°. Accordingly, saddle 706 is sloped at less than 30° and preferably about 20° while saddle 708 is sloped at about 30° and preferably more than 30°.

The implant 760 as shown in Fig. 78 is similar to implant 700 in Fig. 74 and is similarly numbered. Implant 760 includes third and fourth legs 714, 716 which have sloping portions 762, 764 which slope toward ends 766, 768 of third and fourth arm 714, 716 respectively. The sloping portions accommodate the form of the lower vertebra against which they are positioned. In the preferred embodiment, the sloping portions are of about 30°. However, it is to be understood that sloping portions which are substantially greater and substantially less than 30° can be included and be within the spirit and scope of the invention.

25

Embodiment of Fig. 79, 80, 80a, 81, 82, 83, 83a, 84, 85, 86 and 87

Another embodiment of the invention is shown in Figs. 79-87 and includes implant 800 (Fig. 86). Implant 800 includes a distracting unit 802 which is shown in left side, plan, and right side views of Figs. 79, 80

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and 81. A perspective view of the distraction unit is shown in Fig. 84. The distracting unit as can be seen in Fig. 80 includes a distracting body 804, with longitudinal axis 805, which body 804 has a groove 806 and a rounded or bulbous end 808 which assist in the placement of the distracting body between adjacent spinous process so that an appropriate amount of distraction can be accomplished. Extending from the distracting body 804 is a first wing 810 which in Fig. 80 is substantially perpendicular to the distracting body 804. Such wings which are not perpendicular to the body are within the spirit and scope of the invention. First wing 810 includes a upper portion 812 and a lower portion 814. The upper portion 810 (Figs. 79) includes a rounded end 816 and a small recess 818. The rounded end 816 and the small recess 818 in the preferred embodiment are designed to accommodate the anatomical form or contour of the L4 (for a L4-L5 placement) or L5 (for a L5-S1 placement) superior lamina of the vertebra. It is to be understood that the same shape or variations of this shape can be used to accommodate other lamina of any vertebra. The lower portion 814 is also rounded in order to accommodate in the preferred embodiment in order to accommodate the vertebrae. The distracting unit further includes a threaded bore 820 which in this embodiment accepts a set screw 822 (Fig. 86) in order to hold a second wing 824 (Figs. 82, 83) in position as will be discussed hereinbelow.

The threaded bore 820 in this embodiment slopes at approximately 45° angle and intersects the slot 806. With the second wing 824 in position, the set screw 822 when it is positioned in the threaded bore 820 can engage and hold the second wing 824 in position in the slot 806.

Turning to Figs. 82, 83 and 85, left side, plan and perspective views of the second wing 824 are depicted. The second wing 824 is

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similar in design to the first wing. The second wing includes an upper portion 826 and a lower portion 828. The upper portion includes a rounded end 830 and a small recess 832. In addition, the second wing 824 includes a slot 834 which mates with the slot 806 of the distracting unit 802. The second wing 824 is the retaining unit of the present embodiment.

As can be seen in Fig. 83 and 86, the second wing or retaining unit 824 includes the upper portion 826 having a first width "a" and the lower portion 828 having a second width "b". In the preferred embodiment, the second width "b" is larger than first width "a" due to the anatomical form or contour of the L4-L5 or L5-S1 laminae. As can be seen in Fig. 83a in second wing or retaining unit 824, the widths "a" and "b" would be increased in order to, as described hereinbelow, accommodate spinous processes and other anatomical forms or contours which are of different dimensions. Further, as appropriate, width "a" can be larger than width "b". Thus, as will be described more fully hereinbelow, the implant can include a universally-shaped distracting unit 802 with a plurality of retaining units 824, with each of the retaining units having different widths "a" and "b". During surgery, the appropriately sized retaining unit 824, width with the appropriate dimensions "a" and "b" can be selected to match to the anatomical form of the patient.

Fig. 86 depicts an assembled implant 800 positioned adjacent to upper and lower laminae 836, 838 (which are shown in dotted lines) of the upper and lower vertebrae. The vertebrae 836, 838 are essentially below the implant 800 as shown in Fig. 86. Extending upwardly from the vertebrae 836, 838, and between the first and second wings 810, 824, are the upper and lower spinous processes 840, 842. It is to be understood that in a preferred embodiment, the fit of the implant

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between the spinous processes can be such that the wings do not touch the spinous processes, as shown in Fig. 86, and be within the spirit and scope of the invention.

The implant 800 includes, as assembled, an upper saddle 844 and the lower saddle 846. The upper saddle 844 has an upper width identified by the dimension "UW". The lower saddle 846 has a lower width identified by the dimension "LW". In a preferred embodiment, the upper width is greater than the lower width. In other embodiments, the "UW" can be smaller than the "LW" depending on the anatomical requirements. The height between the upper and lower saddles 844, 846 is identified by the letter "h". These dimensions are carried over into Fig. 87 which is a schematic representation of the substantially trapezoidal shape which is formed between the upper and lower saddles. The table below gives sets of dimensions for the upper width, lower width, and height as shown in Fig. 87. This table includes dimensions for some variations of this embodiment.

TABLE

| <i>Variation</i> | <i>1</i> | <i>2</i> | <i>3</i> |
|-------------------------|-----------------|-----------------|-----------------|
| Upper Width | 8 | 7 | 6 |
| Lower Width | 7 | 6 | 5 |
| Height | 10 | 9 | 8 |

For the above table, all dimensions are given in millimeters.

For purposes of surgical implantation of the implant 800 into a patient, the patient is preferably positioned on his side (arrow 841 points up from an operating table) and placed in a flexed (tucked) position in order to distract the upper and lower vertebrae.

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In a preferred procedure, a small incision is made on the midline of the spinous processes. The spinous processes are spread apart or distracted with a spreader. The incision is spread downwardly toward the table, and the distracting unit 802 is preferably inserted upwardly
5 between the spinous processes 840 and 842 in a manner that maintains the distraction of spinous processes. The distracting unit 802 is urged upwardly until the distracting or bulbous end 808 and the slot 806 are visible on the other wide of the spinous process. Once this is visible, the incision is spread upwardly away from the table and the retaining unit or
10 second wing 824 is inserted into the slot 806 and the screw 822 is used to secure the second wing in position. After this had occurred, the incisions can be closed.

An alternative surgical approach requires that small incisions be made on either side of the space located between the spinous
15 processes. The spinous processes are spread apart or distracted using a spreader placed through the upper incision. From the lower incision, the distracting unit 802 is preferably inserted upwardly between the spinous processes 840 and 842 in a manner that urges the spinous processes apart. The distracting unit 802 is urged upwardly until the
20 distracting or bulbous end 808 and the slot 806 are visible through the second small incision in the patient's back. Once this is visible, the retaining unit or second wing 824 is inserted into the slot 806 and the screw 822 is used to secure the second wing in position. After this has occurred, the incisions can be closed.

25 The advantage of either of the above present surgical procedures is that a surgeon is able to observe the entire operation, where he can look directly down onto the spinous processes as opposed to having to view the procedure from positions which are to the right and to the left of the spinous processes. Generally, the incision is as small as possible

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and the surgeon is working in a bloody and slippery environment. Thus, an implant that can be positioned directly in front of a surgeon is easier to insert and assemble than an implant which requires the surgeon to shift from side to side. Accordingly, a top-down approach, as an approach along a position to anterior line is preferred so that all aspects of the implantation procedure are fully visible to the surgeon at all times. This aides in the efficient location of (i) the distracting unit between the spinous processes, (ii) the retaining unit in the distracting unit, and (iii) finally the set screw in the distracting unit.

Fig. 80a shows an alternative embodiment of the distracting unit 802a. This distracting unit 802a is similar to distracting unit 802 in Fig. 80 with the exception that the bulbous end 808a is removable from the rest of the distracting body 804a as it is screwed into the threaded bore 809. The bulbous end 808a is removed once the distracting unit 802a is positioned in the patient in accordance with the description associated with Fig. 86. The bulbous end 808a can extend past the threaded bore 820 by about 1 cm in a preferred embodiment.

Embodiment of Figs. 88, 89, 90 and 91

Another embodiment of the invention is shown in Figs. 88, 89, 90 and 91. In this embodiment, the implant is identified by the number 900. Other elements of implant 900 which are similar to implant 800 are similarly numbered but in the 900 series. For example, the distracting unit is identified by the number 902 and this is in parallel with the distracting unit 802 of the implant 800. The distracting body is identified by the number 904 in parallel with the distracting body 804 of the implant 800. Focusing on Fig. 90, the distracting unit 902 is depicted in a perspective view. The distracting unit includes slot 906 which is wider at the top than at the bottom. The reason for this is that the wider upper

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portion of the slot 906, which is wider than the second wing 924 (Fig. 89), is used to allow the surgeon to easily place the second wing 924 into the slot 906 and allow the wedge-shaped slot 906 to guide the second wing 924 to its final resting position. As can be see in Fig. 91,
5 in the final resting position, the largest portion of the slot 906 is not completely filled by the second wing 924.

The end 908 of implant 900 is different in that it is more pointed, having sides 909 and 911 which are provided at about 45° angles (other angles, such as by way of example only, from about 30° to about 60°
10 are within the spirit of the invention), with a small flat tip 913 so that the body 904 can be more easily urged between the spinous processes.

The distracting unit 902 further includes a tongue-shaped recess 919 which extends from the slot 906. Located in the tongue-shaped recess is a threaded bore 920.

15 As can be seen in Fig. 89, a second wing 924 includes a tongue 948 which extends substantially perpendicular thereto and between the upper and lower portions 926, 928. The tab 948 includes a bore 950. With the second wing 924 positioned in the slot 906 of the distracting unit 902 and tab 948 positioned in recess 919, a threaded set screw 922
20 can be positioned through the bore 950 and engage the threaded bore 920 in order to secure the second wing or retaining unit 924 to the distracting unit 902. The embodiment 900 is implanted in the same manner as embodiment 800 previously described. In addition, as the bore 922 is substantially perpendicular to the distracting body 904 (and
25 not provided at an acute angle thereto), the surgeon can even more easily secure the screw in place from a position directly behind the spinous processes.

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Embodiment of Figs. 92, 92a, 92b, 93, 93a, 93b, 93c, 93d, 94, 94a, 94b, 95, 95a, and 96

Still a further embodiment of the invention is depicted in Figs. 92, and 92a. In this embodiment, the implant 1000 as can be seen in Fig. 5 92a includes a central elongated body 1002 which has positioned at one end thereof a first wing 1004. Wing 1004 is similar to the first wing previously described with respect to the embodiment of Fig. 88. Bolt 1006 secures wing 1004 to body 1002 in this embodiment. Bolt 1006 is received in a bore of the body 1002 which is along the longitudinal 10 axis 1008 of body. It is to be understood that in this embodiment, the first unit is defined by the central body 1002, the first wing 1004, and the guide 1010.

Alternatively, the first wing can be secured to the central body with a press fit and detent arrangement as seen in Fig. 93c. In this 15 arrangement, the first wing has a protrusion 1040 extending preferably about perpendicularly from the first wing, with a flexible catch 1042. The protrusion and flexible catch are press fit into a bore 1044 of the central body with the catch received in a detent 1046.

In yet another alternative embodiment, the first wing can be 20 designed as shown in Fig. 93d with the protrusion directed substantially parallel to the first wing from a member that joins the first wing to the protrusion. Thus in this embodiment, the first wing is inserted into the body along the same direction as the second wing is inserted.

Positioned at the other end of the central body 1002 is a guide 25 1010. In this particular embodiment, guide 1010 is essentially triangularly-shaped so as to be a pointed and arrow-shaped guide. Alternatively, guide 1010 could be in the shape of a cone with lateral truncated sides along the longitudinal axis 1008. Guide 1010 includes

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a recess 1012 having a threaded bore 1014. Recess 1012 is for receiving a second wing 1032 as will be described hereinbelow.

5 Additionally, it is also to be understood that the guide 1010 can be bulbous, cone-shaped, pointed, arrow-shaped, and the like, in order to assist in the insertion of the implant 1000 between adjacent spinous processes. It is advantageous that the insertion technique disturb as little of the bone and surrounding tissue or ligaments as possible in order to (1) reduce trauma to the site and facilitate early healing, and (2) not destabilize the normal anatomy. It is to be noted that with the
10 present embodiment, there is no requirement to remove any of the bone of the spinous processes and depending on the anatomy of the patient, there may be no requirement to remove or sever ligaments and tissues immediately associated with the spinous processes.

15 The implant 1000 further includes a sleeve 1016 which fits around and is at least partially spaced from the central body 1002. As will be explained in greater detail below, while the implant may be comprised of a bio-compatible material such as titanium, the sleeve is comprised preferably of a super-elastic material which is by way of example only, a nickel titanium material (NiTi), which has properties which allow it to
20 withstand repeated deflection without fatigue, while returning to its original shape. The sleeve could be made of other materials, such as for example titanium, but these materials do not have the advantages of a super-elastic material.

25 Fig. 93a is a cross-section through the implant 1000 depicting the central body 1002 and the sleeve 1016. As can be seen from the cross-section of Fig. 93a in a preferred embodiment, both the central body 1002 and the sleeve 1016 are substantially cylindrical and oval or ecliptically-shaped. An oval or elliptical shape allows more of the spinous process to be supported by the sleeve, thereby distributing the

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load between the bone and the sleeve more evenly. This reduces the possibility of fracture to the bone or bone resorption. Additionally, an oval or elliptical shape enhances the flexibility of the sleeve as the major axis of the sleeve, as described below, is parallel to the longitudinal direction of the spinous process. However, other shapes such as round cross-sections can come within the spirit and scope of the invention.

In this particular embodiment, the central body 1002 includes elongated grooves 1018, along axis 1008, which receives elongated spokes 1020 extending from the internal surface of the cylinder 1016.

In a preferred embodiment, both the cross-section of the central body and the sleeve have a major dimension along axis 1022 and a minor dimension along axis 1024 (Fig. 93a). The spokes 1020 are along the major dimension so that along the minor dimension, the sleeve 1016 can have its maximum inflection relative to the central body 1002. It is to be understood that the central body along the minor dimension 1024 can have multiple sizes and can, for example, be reduced in thickness in order to increase the ability of the sleeve 1016 to be deflected in the direction of the central body 1002.

Alternatively as can be seen in Fig. 93b, the central body 1002 can include the spokes 1020 and the sleeve 1016 can be designed to include the grooves 1018 in order to appropriately space the sleeve 1016 from the central body 1002.

In other embodiments, the sleeve can have minor and major dimensions as follows:

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| | <u>Minor Dimension</u> | <u>Major Dimension</u> |
|---|------------------------|------------------------|
| | 6 mm | 10 mm |
| | 8 mm | 10.75 mm |
| | 12 mm | 14 mm |
| 5 | 6 mm | 12.5 mm |
| | 8 mm | 12.5 mm |
| | 10 mm | 12.5 mm |

10 In one preferred embodiment, said sleeve has a cross-section with a major dimension and a minor dimension and said major dimension is greater than said minor dimension and less than about two times said minor dimension. In said embodiment, said guide has a cross-section which is adjacent to said sleeve with a guide major dimension about equal to said sleeve major dimension and a guide
15 minor dimension about equal to said sleeve minor dimension. Further in said embodiment, said guide extends from said central body with a cross-section which reduces in size in a direction away from said central body.

20 In another preferred embodiment, said guide is cone-shaped with a base located adjacent to said sleeve. Further, said guide has a base cross-section about the same as the oval cross-section of said sleeve.

Thus, from the above, it is evident that preferably a major dimension of the sleeve correspond with a major dimension of the central body and a minor dimension of the sleeve corresponds with a
25 minor dimension of the central body. Additionally, it is evident that the major dimension of the sleeve 1016 is substantially perpendicular to a major dimension of the first wing 1004 along longitudinal axis 1030 (Fig. 92a). This is so that as discussed above, when the implant 1000 is properly positioned between the spinous processes, a major portion of
30 the sleeve comes in contact with both the upper and lower spinous

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processes in order to distribute the load of the spinous processes on the sleeve 1016 during spinal column extension.

As indicated above, the preferred material for the sleeve 1016 is a super-elastic material and more preferably one comprised of an alloy of nickel and titanium. Such materials are available under the trademark Nitinol. Other super-elastic materials can be used as long as they are bio-compatible and have the same general characteristics of super-elastic materials. In this particular embodiment, a preferred super-elastic material is made up of the following composition of nickel, titanium, carbon, and other materials as follows:

| | |
|----------|------------------|
| Nickel | 55.80% by weight |
| Titanium | 44.07% by weight |
| Carbon | <0.5% by weight |
| Oxygen | <0.5% by weight |

In particular, this composition of materials is able to absorb about 8% recoverable strain. Of course, other materials which can absorb greater and less than 8% can come within the spirit and scope of the invention. This material can be repeatably deflected toward the central body and returned to about its original shape without fatigue. Preferably and additionally, this material can withstand the threshold stress with only a small amount of initial deforming strain and above the threshold stress exhibit substantial and about instantaneous deformation strain which is many times the small amount of initial deforming strain. Such a characteristic is demonstrated in Fig. 118 where it is shown that above a certain threshold stress level, deformation strain is substantially instantaneous up to about 8%. Fig. 118 shows a loading and unloading curve between stress and deformation strain for a typical type of super-elastic material as described above.

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Preferably, the above super-elastic material is selected to allow deformation of up to about, by way of example only, 8%, at about 20 lbs. to 50 lbs. force applied between a spinous processes. This would cause a sleeve to deflect toward the central body absorbing a substantial amount of the force of the spinous processes in extension. Ideally, the sleeves are designed to absorb 20 lbs. to 100 lbs. before exhibiting the super-elastic effect (threshold stress level) described above. Further, it is possible, depending on the application of the sleeve and the anatomy of the spinal column and the pairs of spinous processes for a particular individual, that the sleeve can be designed for a preferable range of 20 lbs. to 500 lbs. of force before the threshold stress level is reached. Experimental results indicate that with spinous processes of an older individual, that at about 400 pounds force, the spinous process may fracture. Further, such experimental results also indicate that with at least 100 pounds force, the spinous process may experience some compression. Accordingly, ideally the super-elastic material is designed to deform or flex at less than 100 pounds force.

In a preferred embodiment, the wall thickness of the sleeve is about 1 mm or 40/1000 of an inch (.040 in.). Preferably the sleeve is designed to experience a combined 1 mm deflection. The combined 1 mm deflection means that there is $\frac{1}{2}$ mm of deflection at the top of the minor dimension and a $\frac{1}{2}$ mm deflection at the bottom of the minor dimension. Both deflections are toward the central body.

In a particular embodiment where the sleeve is more circular in cross-section, with an outer dimension of 0.622 in. and a wall thickness of 0.034 in., a 20 lb. load causes a 0.005 in. deflection and a 60 lb. load causes a 0.020 in. deflection (approximately $\frac{1}{2}$ mm). A 100 lb. load would cause a deflection of about 0.04 in. or approximately 1 mm.

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Thus in summary, the above preferred super-elastic material means that the sleeve can be repeatedly deflected and returned to about its original shape without showing fatigue. The sleeve can withstand a threshold stress with a small amount of deforming strain and at about said threshold stress exhibit about substantially instantaneous deformation strain which is many times the small amount of the forming strain. In other words, such super-elastic qualities mean that the material experiences a plateau stress where the material supports a constant force (stress) over very large strain range as exhibited in Fig. 118.

It is to be understood that for this particular embodiment, bar stock of the super-elastic material is machined into the appropriate form and then heat treated to a final temperature to set the shape of the material by increasing the temperature of the material to 932° Fahrenheit and holding that temperature for five (5) minutes and then quickly quenching the sleeve in water. It is also to be understood that preferably the present nickel titanium super-elastic alloy is selected to have a transition temperature A_f of about 59° Fahrenheit (15°C). Generally for such devices the transition temperature can be between 15°C to 65°C (59°F to 149°F), and more preferably 10°C to 40°C (50°F to 104°F). Preferably, the material is maintained in the body above the transition temperature in order to exhibit optimal elasticity qualities.

Alternatively, and preferably, the sleeve can be fabricated by wire Electrical Discharge Machining (EDM) rather than machined. Additionally, the sleeve can be finished using a shot blast technique in order to increase the surface strength and elasticity of the sleeve.

Top and side views of the second wing 1032 are shown in Figs. 94 and 95. Second wing 1032 as in several past embodiments includes a tab 1034 with a bore 1036 which aligns with the bore 1014 of the

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guide 1010. In this particular embodiment, the second wing 1032 includes a cut-out 1038 which is sized to fit over the guide 1010, with the tab 1034 resting in the recess 1012 of the guide 1010.

5 An alternative configuration of the second wing 1032 is depicted in Fig. 94a. In this configuration, the second wing 1032 is held at acute angle with respect to the tab 1034. This is different from the situation in the embodiment of Figs. 94 and 95 where the second wing is substantially perpendicular to the tab. For the embodiment of the second wing in Fig. 94a, such embodiment will be utilized as appropriate
10 depending on the shape of the spinous processes.

With respect to the alternative second wing 1032 depicted in Figs. 94b and 95a, elongated tab 1034 has a plurality of closely positioned bores 1036. The bores, so positioned, appear to form a scallop shape. Each individual scallop portion of the bore 1036 can selectively hold the
15 bolt in order to effectively position the second wing 1032 in three different positions relative to the first wing 1004. The cut-out 1038 (Fig. 95a of this alternative embodiment) is enlarged over that of Fig. 95 as in a position closest to the first wing 1004, the second wing 1032 is immediately adjacent and must conform to the shape of the sleeve
20 1016.

Embodiment of Fig. 97

Implant 1050 of Fig. 97 is similar to the implant 1000 in Fig. 92 with the major difference being that a second wing is not required. The
25 implant 1050 includes a central body as does implant 1000. The central body is surrounded by a sleeve 1016 which extends between a first wing 1004 and a guide 1010. The guide 1010 in this embodiment is substantially cone-shaped without any flats and with no bore as there is no need to receive a second wing. The sleeve and the central body as

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well as the first wing and guide act in a manner similar to those parts of the implant 1000 in Fig. 92. It is to be understood a cross-section of this implant 1050 through sleeve 1016 can preferably be like Fig. 93a. This particular embodiment would be utilized in a situation where it was
5 deemed impractical or unnecessary to use a second wing. This embodiment has the significant advantages of the sleeve being comprised of super-elastic alloy materials as well as the guide being utilized to guide the implant between spinous processes while minimizing damage to the ligament and tissue structures found around
10 the spinous processes.

Embodiment of Fig. 98

Implant 1060 is depicted in Fig. 98. This implant is similar to the implants 1000 of Fig. 92 and the implant 1050 of Fig. 97, except that this
15 implant does not have either first or second wings. Implant 1060 includes a sleeve 1016 which surrounds a central body just as central body 1002 of implant 1000 in Fig. 93. It is to be understood that a cross-section of this implant 1060 through sleeve 1016 can preferably be like Fig. 93a. Implant 1060 includes a guide 1010 which in this
20 preferred embodiment is cone-shaped. Guide 1010 is located at one end of the central body. At the other end is a stop 1062. Stop 1062 is used to contain the other end of the sleeve 1016 relative to the central body. This embodiment is held together with a bolt such as bolt 1006 of Fig. 93 that is used for the immediate above two implants. For the
25 implant 1060 of Fig. 98, such a device would be appropriate where the anatomy between the spinous processes was such that it would be undesirable to use either a first or second wing. However, this embodiment affords all the advantageous described hereinabove (Figs.

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92 and 97) with respect to the guide and also with respect to the dynamics of the sleeve.

Embodiment of Figs. 99 and 100

5 Figs. 99 and 100 depict an implant system 1070. Implant system 1070 includes a sleeve 1072 which is similar to and has the advantageous of sleeve 1016 of the embodiment in Fig. 92. Sleeve 1072 does not, however, have any spokes. Additionally, implant system 1070 includes an insertion tool 1074. Insertion tool 1074 includes a
10 guide 1076 which in a preferred embodiment is substantially cone-shaped. Guide 1076 guides the insertion of the sleeve 1072 and the insertion tool 1074 between adjacent spinous processes. The insertion tool 1074 further includes a central body 1078, a stop 1080, and a handle 1082. The guide 1076 at its base has dimensions which are
15 slightly less than the internal dimensions of the sleeve 1074 so that the sleeve can fit over the guide 1076 and rest against the stop 1080. The tool 1074 with the guide 1076 is used to separate tissues and ligaments and to urge the sleeve 1072 in the space between the spinous processes. Once positioned, the guide insertion tool 1074 can be
20 removed leaving the sleeve 1072 in place. If desired, after the sleeve is positioned, position maintaining mechanisms such as springy wires 1084 made out of appropriate material such as the super-elastic alloys and other materials including titanium, can be inserted using a cannula through the center of the sleeve 1072. Once inserted, the ends of the
25 retaining wires 1084 (Fig. 99) extend out of both ends of the sleeve 1072, and due to this springy nature, bent at an angle with respect to the longitudinal axis of the sleeve 1072. These wires help maintain the position of the sleeve relative to the spinous processes.

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Embodiment of Figs. 101, 102, 102a, 103, 104, 105, 106, and 107

Another embodiment of the invention can be seen in Fig. 101 which includes implant 1100. Implant 1100 has many similar features that are exhibited with respect to implant 1000 in Fig. 92. Accordingly, elements with similar features and functions would be similarly
5 numbered. Additionally, features that are different from implant 1100 can be, if desired, imported into and become a part of the implant 1000 of Fig. 92.

As with implant 1000, implant 1100 includes a central body 1002
10 (Fig. 102) with a first wing 1004 and a bolt 1006 which holds the first wing and the central body together. In this particular embodiment, the central body is made in two portions. The first portion 1102 is in the shape of a truncated cone with an oval or elliptical base and a second portion 1104 includes a cylindrical central portion with a distal end in the
15 shape of a truncated cone 1103 with an oval or elliptical base. In addition, in this particular embodiment, formed with the central body is the guide 1010 which has an oval or elliptical base. Bolt 1006 is used to secure the first wing through the second portion 1104 with the first portion 1102 held in-between. In this particular embodiment, the guide
20 1010 in addition to including recess 1012 and bore 1014 includes a groove 1106 which receives a portion of the second wing 1032.

In this particular embodiment, the sleeve 1016 is preferably oval or elliptical in shape as can be seen in Fig. 102a. The central body can be oval, elliptical or circular in cross-section, although other shapes are
25 within the spirit and scope of the invention. The sleeve 1016 held in position due to the fact that the truncated conical portion 1102 and the corresponding truncated conical portion 1103 each have a base that is elliptical or oval in shape. Thus, the sleeve is held in position so that preferably the major dimension of the elliptical sleeve is substantially

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perpendicular to the major dimension of the first wing. It is to be understood that if the first wing is meant to be put beside the vertebrae so that the first wing is set at an angle other than perpendicular with respect to the vertebrae and that the sleeve may be held in a position
5 so that the major dimension of the sleeve is at an angle other than perpendicular to the major dimension of the first wing and be within the spirit and scope of the invention. This could be accomplished by tightening bolt 1006 with the first wing 1004 and sleeve 1016 so positioned. In such a configuration, the major dimension of the sleeve
10 would be preferably positioned so that it is essentially parallel to the length of the adjacent spinous processes. So configured, the elliptical or oval shape sleeve would bear and distribute the load more evenly over more of its surface.

It is to be understood that the sleeve in this embodiment has all
15 the characteristics and advantages described hereinabove with respect to the above-referenced super-elastic sleeves.

The second wing as discussed above, can come in a variety of shapes in order to provide for variations in the anatomical form of the spinous processes. Such shapes are depicted in Figs. 103, 104, 105,
20 106, and 107. In each configuration, the second wing 1032 has an upper portion 1108 and a lower portion 1110. In Fig. 104, the lower portion is thicker than the upper portion in order to accommodate the spinous process, where the lower spinous process is thinner than the upper spinous process. In Fig. 105, both the upper and lower portions are
25 enlarged over the upper and lower portions of Fig. 103 to accommodate both the upper and lower spinous processes being smaller. That is to say that the space between the upper and lower portions of the first and second wings are reduced due to the enlarged upper and lower portions of the second wing.

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Alternative embodiments of second wings, as shown in Figs. 104 and 105, are depicted in Figs. 106 and 107. In these Figs. 106 and 107, the second wing 1032 accommodates the same anatomical shape and size of the spinous processes as does the second wing in Figs. 104 and 105 respectively. However, in the embodiments of the second wing 1032 of Figs. 106 and 107, substantial masses have been removed from the wings. The upper and lower portions 1108 and 1110 are essentially formed or bent in order to extend from the central portion 1112 of the second wing 1032.

It is to be understood that in this embodiment, if desired, the second wing may not have to be used, depending on the anatomy of the spinal column of the body, and this embodiment still has the significant advantages attributable to the guide 1010 and the functionality of the sleeve 1016.

Embodiment of Figs. 108, 109, and 110

The implant 1120 as shown in Figs. 108 and 109, is similar to implant 1100 which is in turn similar to implant 1000. Such similar details have already been described above and reference here is made to the unique orientation of the first and second wings 1122 and 1124. These wings have longitudinal axis 1126 and 1128 respectfully. As can be seen in these figures, the first and second wings 1122, 1124 have been rotated so that they both slope inwardly and if they were to continue out of the page of the drawing of Fig. 108, they would meet to form an A-frame structure as is evident from the end view of Fig. 109. In this particular embodiment, as can be seen in Figs. 109 and 110, the tab 1034 is provided an acute angle to the remainder of the second wing 1124. Further, the groove 1018 formed in the implant is sloped in order to accept the second wing 1124. Accordingly, this present implant 1120

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is particularly suited for an application where the spinous process is wider adjacent to the vertebral body and then narrows in size at least some distance distally from the vertebral body. It is to be understood that a cross-section of this implant 1120 through sleeve 1016 can preferably be like Fig. 93a.

Embodiment of Figs. 111, 112, 113, 114, 115, 116, and 117

An additional embodiment of the implant 1150 is shown in Fig. 111. Implant 1150 has features similar to those described with respect to Fig. 94b.

Implant 1150 includes a central body 1152 with a first wing 1154, where central body 1152 includes elongated groove 1156 which extends to the guide 1158. A screw 1160 is received in a threaded bore located in the elongated groove 1156.

The second wing 1162 includes a central body 1164 which is substantially perpendicular to the second wing 1162.

The central body 1164 includes a plurality of bores 1166 provided therein. These bores are formed adjacent to each other in order to define a plurality of scallops, each scallop capable of retaining bolt 1160 therein. As can be seen in Fig. 114, the second wing includes a cut-out 1168 such that with the central body 1164 of the second wing received in the groove 1156 of the central body associated with the first wing, the remainder of the second wing is received over the central body 1152 of the implant 1150. With this implant 1150, the distance between the first and second wings can be adjusted by selectively placing the bolt 1160 through one of the five specified bores defined by the scalloped plurality of bores 1166. Accordingly, Fig. 112 depicts the implant where the first and second wings are widest apart in order to accommodate spinous processes of greater thickness. Fig. 111 shows the middle position

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between the first and second wings in order to accommodate average size spinous processes.

It is to be understood that preferably during the surgical process, the central body 1152 is urged between spinous processes. After this
5 has occurred, the second wing is guided by the other sides of the spinous processes from a path which causes the plane of the second wing to move substantially parallel to the plane of the first wing until the central body 1164 associated with the second wing 1162 is received in the groove of 1156 of the central body 1152 associated with the first
10 wing 1154. After this has occurred, the bolt 1160 is positioned through aligned bores associated with the second wing 1162 and the central body 1152 in order to secure the second wing to the central body.

While embodiment 1150 does not depict a sleeve such as sleeve 1016, such a sleeve 1016 could be placed over body 1152 and be within
15 the spirit of the invention.

Embodiments of Figs. 119a, 119b, 120a, 120b, 121a, 121b, 122a, 122b, 122c, 123a, 123b, 124a, 124b, and 124c

Implant 1200 of the invention is depicted in Figs. 119a and 119b.
20 This implant includes the first wing 1202 and sleeve 1204 and a guide 1206. An alternative to this embodiment further includes, as required, second wing 1208 as depicted in Figs. 120a and 120b.

As can be seen in Fig. 121a and 121b, the first wing 1202 includes a bore which receives a central body 1210. Preferably, the
25 central body is pressed fit through the bore of the first wing although it is to be understood that other securing mechanisms such as through the use of threads and still other mechanisms can be used to accomplish this task. Additionally, in this particular embodiment first and second pins 1212 extend from the first wing 1202, each along an axis which is

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substantially parallel to the longitudinal axis 1214 of the central body 1210. In this particular embodiment, the distal end 1216 of the central body 1210 is threaded in order to be coupled to the guide 1206.

As can be seen in Figs. 122a, 122b and 122c, the guide 1206 in
5 this particular embodiment is pointed in order to allow the implant to be inserted between, and if necessary distract, adjacent spinous processes. The guide 206 includes a threaded bore 1218 which is designed to accept the threaded end 1216 of the central body 1210 in order to secure the guide to the central body and additionally for
10 purposes of retaining the sleeve between the guide 1206 and the first wing 1202.

As can be seen in Fig. 123a the sleeve 1204 is preferably cylindrical, and oval or elliptical in shape in cross-section. It is to be understood that sleeve 1204 can have other shapes as described
15 throughout the specification and be within the spirit and scope of the invention. In this particular embodiment, sleeve 1204 has at least one major diameter and one minor diameter in cross-section. Sleeve 1204 includes a central bore 1220 which extends the length of sleeve 1204 and curve grooves 1222 which are formed about central bore 1220 and
20 extend only part way into the body of the sleeve. In this particular embodiment, the curved grooves 1222 describe an arc of about 60°. It is to be understood that in other embodiment, this arc can be less than 60° and extend past 120°.

The sleeve 1204 is received over the central body 1210 of the
25 implant 1200 and can rotate thereon about the longitudinal axis 1214 of the central body 1210. When this particular embodiment is assembled, the grooves 1222 have received therein the pins 1212 that extend from the first wing 1202. Accordingly, the pins inserted in the grooves 1222 assist in the positioning of the sleeve relative to the remainder of the

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implant 1200. With the pins 1212 received in the curved grooves 1222, the pins limit the extent of the rotation of the sleeve about the central body and relative to the first wing.

5 As can be seen in Figs. 124a, 124b, and 124c, the sleeve is free to rotate relative to the longitudinal axis of the central body 1210 and thus relative to the first wing 1202 of the embodiment shown in Figs. 119a and 119b. The sleeve can rotate relative to a second wing 1208, when the second wing is utilized in conjunction with the embodiment of Figs. 119a and 119b. The pins limit the rotation of the sleeve. In an
10 alternative embodiment, the pins are eliminated so that the sleeve can rotate to any position relative to the first wing.

It is to be understood that the sleeve can be comprised of biologically acceptable material such as titanium. Additionally, it can be comprised of super-elastic material such as an alloy of nickel and
15 titanium, much as described hereinabove with respect to other embodiments.

The great advantage of the use of the sleeve 1204 as depicted in the embodiment of Figs. 119a and 119b is that the sleeve can be rotated and repositioned with respect to the first wing 1202, and/or the
20 second wing 1208 should the second wing be used in the embodiment, in order to more optimally position the implant 1200 between spinous processes. It is to be understood that the cortical bone or the outer shell of the spinous processes is stronger at an anterior position adjacent to the vertebral bodies of the vertebra than at a posterior position distally
25 located from the vertebral bodies. Accordingly, there is some advantage of having the implant 1200 placed as close to the vertebral bodies as is possible. In order to facilitate this and to accommodate the anatomical form of the bone structures, as the implant is inserted between the vertebral bodies and urged toward the vertebral bodies, the sleeve 1204

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can be rotated relative to the wings, such as wing 1202, so that the sleeve is optimally positioned between the spinous processes, and the wing 1202 is optimally positioned relative to the spinous processes. Without this capability, depending on the anatomical form of the bones,
5 it is possible for the wings to become somewhat less than optimally positioned relative to the spinous processes.

Embodiments of Figs. 125, 126, and 127

Figs. 125, 126 and 127 depict three alternative embodiments of
10 the invention as can be seen through a line parallel to line 124-124 of Fig. 119b.

In Fig. 125, the sleeve 1204 is rotatable about central body 1210. In this embodiment, however, the sleeve 1204 design does not include the grooves 1222 as previously depicted in the embodiment shown in
15 Fig. 123a. Thus, without pins, the sleeve is completely free to rotate about the central body 1210.

An alternative embodiment is shown in Fig. 126. In this embodiment, the sleeve 1204 is essentially a thin wall cylinder which is spaced from the central body 1210. Sleeve 1204 is free to move
20 relative to central body 1210. Sleeve 1204 can rotate relative to central body 1210. In addition, sleeve 1204 can take a somewhat cocked or skewed position relative to central body 1210.

A further embodiment, it is shown in Fig. 127. This embodiment is somewhat similar to the embodiment shown in Fig. 126 except that in
25 this case, several pins project from the first wing in order to somewhat limit and restrict the motion of the sleeve 1204. As shown in Fig. 127, four pins are depicted. It is to be understood however that such an embodiment can include one, two, three, four or more pins and be within the spirit and scope of the invention. It is to be understood that if the

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embodiment is used with a second wing, that similar pins can extend from the second wing. However, in the embodiment using a second wing, the pins would preferably be somewhat flexible so that they could snap into the inside of the sleeve 1204 as the second wing is inserted relative to the central body and secured in place. In the embodiment shown in Fig. 127, the sleeve 1204 is free to rotate about the longitudinal axis of the central body 1210 and is somewhat restricted in this motion and its ability to become skewed relative to the longitudinal axis of the central body by the pins.

Embodiments of Figs. 128 and 129

The embodiments of Fig. 128 is an advantageous alternative to that of Fig. 93a. In this embodiment, the central body 1002 is similar to that as shown in Fig. 93a. The sleeve 116 is comprised of two sleeve portions 1016a and 1016b. The sleeve portions are preferably formed from flat stock material which is substantially easier to form than having the sleeve formed or machined from solid bar stock material. A further advantage of the sleeve 1016, if formed of super-elastic material, is that the sleeve can be formed in a manner which optimizes the super-elastic characteristics of such material in order to enhance its ability to repeatedly deflect under load. In this particular embodiment, the sleeve portions 1016a and 1016b are somewhat C-shaped and then after being formed, are snapped into the grooves of the central body 1002.

An alternative embodiment of the invention is shown in Fig. 128. This embodiment is most favorably used with the embodiment of Fig. 119a and 119b. In this particular embodiment, the sleeve 1204 is designed to rotate about the central body 1210. Sleeve 1204 includes a central member 1230 which includes a bore that receives the central body 1210. The central member 1230 is rotatable about the central

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body 1210 of the implant 1200. The central member 1230 includes first and second grooves 1232 and 1234. These grooves can receive C-shaped sleeve members 1204a and 1204b. These C-shaped sleeve members are similar in construction and design to the C-shaped sleeve members shown above with respect to Fig. 128. These sleeve members can be snapped into position relative to the central member 1230 of the sleeve 1204. It is to be understood that other mechanisms can be used to secure the C-shaped sleeve member relative to the central member of the sleeve and be within the spirit and scope of the invention. Further, it is to be understood that the sleeve members 1204a and 1204b can be formed from a single flat stock material such that one of the grooves 1232 and 1234 receives continuous piece of flat material which has been appropriately bent and the other grooves receives two ends of the sleeve.

15

Embodiments of Figs. 130-136

Embodiment 2000 of the supplemental spine fixation device of the invention is depicted in Fig. 130. This embodiment 2000 includes a hub 2002 to which is adjustably secured a first hook member 2004 and a second hook member 2006. First hook member 2004 includes a hook 2008 which is more fully described hereinbelow, and a shaft 2010 extending therefrom. Similarly, second hook member 2006 includes a hook 2012 and a shaft 2014 extending therefrom. As described more fully hereinbelow, hook 2008 is swivelly or pivotably mounted to shaft 2010. It is to be understood that the description and functionality of first hook member 2004 applies equally well to that of second hook member 2006. The shaft 2010 in this embodiment includes a rack 2016 which can mate selectively with rack 2018 of hook member 2006. These two racks 2016 and 2018 interlock in a multitude of positions in order to

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adjust the position of first and second hook members 2004 and 2006, relative to each other and relative to the hub 2002. The shafts 2010 and 2014 are positioned through bore 2020 in the hub 2002, selectively interlocked together and are then lockingly positioned using a locking mechanism such as the screw 2022. As is described more fully below, the hooks 2008 and 2012 are designed and shaped to fit around spinous processes. Further, the hooks 2008 and 2012 are swivelly mounted to the shafts 2010 and 2014 in order to accommodate the various sizes, shapes, and positions of spinous processes of the human population.

Movably mounted to the hub 2002 is a shaft 2024 (Fig. 131) and extending from the shaft 2024 is an inter-spinous process guide 2026. The shaft 2024 at a proximal end includes a crossbar or tab 2028 which is slidably or movably received in a slot 2030 of the hub 2002. Once the tab 2028 is received in the slot 2030, the slot can be pinched off or slightly deformed at its open end using a punch or other mechanism in order to prevent the tab 2028, and thus the shaft 2024 and the guide 2026 from being removed from the hub 2002. With the tab 2028 located in the slot 2030, the shaft and also the guide 2026 extending from the distal end of the shaft 2024 are free to move relative to the hub and also relative to the first hook member 2004 and the second hook member 2006. This movement, as well as the ability of the hooks 2008 and 2012 to swivel on the shafts 2010 and 2014, allow the embodiment 2000 to conform to the spinous process anatomy.

Movably mounted on the shaft 2024 is a spacer or sleeve 2032. Spacer 2032 includes a central bore 2034 through which the shaft 2024 extends. The spacer 2032 is thus able to rotate about the shaft 2034. The spacer 2032 is cylindrical and in this particular embodiment is oval or elliptical in shape. In addition, the base of the guide 2026 is also

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somewhat elliptical in shape in order to make a smooth transition between the guide 2026 and the spacer 2032 as the guide and spacer are inserted between the spinous processes in order to distract apart the spinous processes during the insertion process. As the spacer 2032 is
5 rotatable on the shaft 2024, and as the spacer 2032 is elliptically shaped, it can be inserted in one position and then as the entire embodiment 2000 is positioned to the final securing position, the spacer 2032 can rotate about the shaft 2024 to accommodate the shape of the space between the spinous processes as the spacer is moved from a
10 posterior position to an anterior position.

The spacer 2032 can include a second alternative spacer embodiment 2036 (Fig. 131a) in substitution for the spacer 2032. Spacer 2036 includes an elongated slot 2038 into which the shaft 2024 can be received. Elongated slot 2038 not only allows the spacer 2036
15 to rotate about the shaft 2024, it also allows it to translate relative to shaft 2024. Such translation in this embodiment is substantially perpendicular to the shaft, in any direction to which the spacer 2036 is rotated. Thus, in this embodiment the degrees of freedom which accommodate the anatomical shape of the spinous processes and the
20 space therebetween, including the ligaments and tissues associated therewith, include (1) the ability of the hooks 2004, 2006 to swivel on the shafts, (2) the ability of the hooks 2004, 2006 to move relative to the hub 2002 and be locked to the hub, (3) the ability of the shaft 2024 to move in the slot 2030 of the hub, and (4) finally the ability of the spacer 2036
25 to both rotate and translate on the shaft 2024.

Before proceeding to more specific details of this embodiment 2000, it is to be understood that the same features of the spacer, the shaft, and the lead-in guide, which are found in other embodiments such as by way of example only, the embodiments of Figs. 10, 16, 20, 22, 86,

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88, 92, and 119b, and other figures, can be incorporated into this embodiment. By way of example only, the implant 2000 can be comprised of stainless steel, titanium or other biologically acceptable materials. The shape of the lead-in plug can be cone shaped, pyramid
5 shaped, and other shapes with a small lead-in cross-section expanding into a larger cross-section which is similar to the cross-section of the spacer 2032, in order to gradually distract apart the spinous processes to a sufficient distance so that the spacer 2032 or the spacer 2036 can conveniently fit between the spinous processes. Further, the spacer, as
10 shown in the other embodiments, can include a spacer made of stainless steel or titanium, or of a super-elastic material or of a silicone. The spacer besides being cylindrical can, from parallel planar end 2040 to parallel planar end 2042, be saddle-shaped along surface 2041 so that the ends are high and the center portions are low in order to more
15 fully accommodate the shape of the spinous processes and also to spread the load across a broader contact surface between the spinous processes and the spacer. For example, the spacer 2032 could have a shape such as the saddle shape defined by the mated together components of the embodiment of Fig. 16. Further, the dimensions of
20 this embodiment as applied to the guide 2006 and the spacer 2032 can be acquired from other embodiments presented herein.

The shape of the guide 2026 and the spacer 2032 is such that for purposes of insertion between spinous processes, the spinous processes to do not need to be altered or cut away in any manner in
25 order to accommodate this implant. Further, the associated ligaments do not need to be cut away and there would be very little or no damage to the other adjacent and surrounding tissues. Similarly, the hook members 2004, 2006, are appropriately shaped and also pivotable so that alteration of the spinous process is not required.

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Returning to Figs. 135a-135f and Fig. 136, the design of the hook members 2004 and 2006 are more fully depicted and described. As indicated above, the description will be made with respect to first hook member 2004. This description applies equally to second hook member 2006. As can be seen in Fig. 135b, the first hook member 2004 includes a shaft 2010 which is received in a bore 2044 of the hook 2008. This bore receives a rounded ball end 2046 seated against a somewhat circular seat 2048. A screw 2050 (Fig. 135f) is received in the bore 2044 in order to retain the rounded ball end 2046. The other end of the bore 2044, end 2052, as can be seen in Fig. 135f is oval or elliptical in shape. This allows the hook 2008 to swivel side to side on the shaft 2010 in order to accommodate the spinous process while somewhat restricting the back and forth rocking of the hook 2008 relative to the shaft 2010. This freedom of motion can be seen in Fig. 136 with respect to the upper spinous processes 2054. The hook can swivel side to side in order to accommodate the shape of the upper spinous processes 2054. The lower hook 2006 additionally can move in order to accommodate the lower spinous processes 2056. As can be seen in Figs. 135c and 135d, the hook 2008 can swivel about 15° on either side of a central longitudinal axis of the shaft 2010.

Additionally with respect to the hook 2008, as can be seen in Figs. 135a, 135b, and 135e, the hook includes a convex inner surface 2058 in order to accommodate the varying surface shape of the spinous processes, and in order to even out the load transferred between the hook and the spinous processes.

The embodiment 2000 can be implanted in a number of methods, preferably, once a spine fixation device is implanted between the vertebral bodies. In this particular embodiment, through a small incision the hub, spacer, and guide are inserted with the guide and spacer

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inserted between the spinous processes. Once this is accomplished, a first hook member and then a second hook member is secured about the respective spinous processes. The shafts of the hook members are then inserted through the bore of the hub 2002 until the spinous processes are brought tight against the spacer. The hooks are appropriately positioned on the spinous processes as depicted in Fig. 136. After this has been accomplished, the securing mechanism 2022 is tightened in order to lock the hooks in place and to secure the spinous processes in a rigid manner relative to each other and relative to the distracting spacer 2032. Alternatively, the spinous ligaments can hold the spinous processes tightly against the spacer and the hooks can be moved and locked into tight contact with the spinous processes.

The above procedure can have variations. By way of example only, the hooks can be inserted first through the incision and then the guide, spacer and hub can be inserted. Once this is accomplished the hooks can be mated to the hub.

In another embodiment and method not depicted, the physician can insert the shaft 2024 on which the spacer 2032 is mounted into the slot 2030 of the hub 2002 and can close off the slot with a securing screw in order to retain shaft 2024. This process is in contrast to the shaft being secured in the slot during the manufacturing process. The securing screw would be similar to securing screw 2022 and would be placed in a bore made at the top of slot 2030. The physician could accordingly insert the tab 2028 of the shaft 2024 in the slot 2030, and then secure the tab in place with the securing screw.

Still an alternative method would be for the device 2000 to be inserted through a larger incision, with device 2000 fully assembled. Once inserted the screw 2022 could be loosened so that the hook members could be positioned around spinous processes at about the

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same time that the guide and spacer are inserted between the spinous processes. Once this is accomplished, the spinous processes could be drawn down tightly around the spacer, with the hooks tightly around the spinous processes and secured firmly into the hub 2002 with the
5 securing screw 2022.

In all of the above procedures, it is advantageous that the device 2000 can address the adjacent spinous processes from one side of the spinous processes and not require exposure of both sides of the spinous processes and thus the procedure is less traumatic to the
10 surgical site.

Still an alternate insertion method would be to insert the device fully assembled with the hook rotated at 90° to the final position shown in Fig. 130. Once the hooks are positioned adjacent to the spinous processes, the hooks could be rotated to the position shown in Fig. 130.
15 Then simultaneously the guide and spacer could be inserted between the spinous processes, as the hooks are positioned about the spinous processes. The hooks are then drawn together, causing the spinous processes to be held firmly against the spacer. Once this is accomplished the screw 2022 can be securely fastened to the hub 2002.

20 With respect to the embodiment of Fig. 130, this embodiment as fully described above can be used as a supplemental fixation or augmentation device for the lumbar level fusion of the L4/L5 vertebrae and above vertebrae, and also for the L5/S1 and below vertebrae. Thus, this device 2000 can be used with respect to fusion of any of the
25 vertebrae up and down the spinous processes.

Embodiments of Figs. 137-140

Another embodiment 2100 of the invention can be seen in Figs. 137-140. Components and features of this embodiment 2100

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which are similar to components and features of the embodiment 2000 have similarly least significant digits. Thus the hub for embodiment 2100 would be 2102. The main difference between the embodiment 2100 and the previously described embodiment 2000 is directed to the
5 hub 2102 and the shafts 2110 and 2114. In this embodiment, the shafts 2110, 2114 are substantially rectangular in cross-sections as opposed to semi-circular as in the previous embodiment of Fig. 130. As can be seen in Fig. 138, shaft 2110 is substantially rectangular in cross-section and include rack or teeth 2116. Shaft 2114 is shaped as a fork with two
10 tines 2115 and 2117. Further, the two tines have rack or teeth 2118. The shaft 2110 of the first hook member 2104 slides between the two tines 2115 and 2117. As can be seen in Fig. 139, with the shaft 2110 slipped between the two tines 2115, 2117 and also with shafts 2110, 2114 located in the rectangular bore 2120 of the hub 2102, the top cap
15 2103 (which is shown both from the top side (Fig. 138) and from the bottom side (Fig. 138b)) can be placed over the hub 2102. The teeth or rack 2105 on the bottom side of the cap 2103, mesh with the teeth or rack 2116 and 2115, 2117 of the first and second hook members. Once this is accomplished, the screw 2122 can be inserted through the
20 indicated bore so that the cap 2103 can tighten down on the hub 2102, locking the shafts 2110 and 2114 of the first and second hook members in place.

All the other features, dimensions, characteristics, materials, methods of insertion, and methods of operation of the embodiment
25 shown in Fig. 138 are similar to or derivations from that shown in the embodiment of Fig. 130.

Embodiments of Figs. 141-143

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Another embodiment of the invention is depicted in Figs. 141-143. This embodiment is similar to the other embodiments 2000 and 2100. This embodiment is numbered 2200. Similar elements, features, methods and aspects have similar numerical designations with respect
5 to the lowest two significant digits. Thus the hub of embodiment 2200 is identified as hub 2202.

In this particular embodiment, the hub has rigidly affixed thereto shaft 2224. Here shaft 2224 does not slide in a slot as happens with respect to the prior two embodiments 200 and 2100. Shaft 2224 can be
10 screwed into hub 2202 or integrally formed with hub 2202. Additionally, the guide 2226 can be integrally formed with the shaft 2224 or in other manners fastened to the shaft 2224 as with a thread mechanism. In this particular embodiment, as can be seen in Fig. 143, the shaft 2224 is integrally formed with the hub 2202 and the shaft 2224 includes a
15 threaded extension 2225 onto which is screwed the guide 2226. For this particular embodiment, the sleeve or spacer 2236 includes the elongated slot 2238 in order to provide for freedom of movement between the sleeve or spacer 2236, the hub 2202, and the first and second hook members 2204 and 2206.

20 In this particular embodiment the shaft 2210 and 2214 are similar to those depicted with respect to the embodiment 2000. In other words each has a rack or teeth which mate with the other. Shafts 2210 and 2214 are inserted through the semi-circular bore 2220 of the hub 2202, and then the cap 2203 is mated on top of the hub 2202. The cap
25 includes a semi-circular bore 2207 which is positioned over the upper shaft 2210. Both bores 2207 and 2220 include ribs, teeth, or threads that run along the length of the bores. These ribs, teeth, or threads are urged against the shafts in order to assist in locking the shafts in place. Alternatively, the ribs, teeth, or threads of the bores can be across the

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length of the bores. The shafts 2210 and 2214 can have teeth, ribs, or threads that are positioned all about the shafts so that the shafts can lock to each other, and so that the teeth, racks or threads on the bores can lock the shafts in place. Once the cap 2203 is positioned over the
5 hub 2202, the screw 2222 is positioned in the bore of the hub 2202 in order to lockingly position the first and second hook members 2204 and 2206 relative to the hub. In particular, with respect to embodiment 2200, the degrees of freedom are attributable to (1) the slot 2238 in the spacer 2236, (2) the shafts 2210 and 2224 which can be positioned relative to
10 each other to position the hooks 2204 and 2206 relative to the hub, and accordingly relative to the spacer, and (3) the ability of the hooks 2204 and 2206 to swivel or pivot.

As indicated above, all the other features, materials, aspects, dimensions, and so forth, of the embodiment 2200 are similar to and can
15 be specified according to the other embodiments 2000 and 2100.

A preferred method of insertion of this embodiment 2200 into a patient is as follows. Initially through a small incision the guide, spacer and hub are inserted so that the guide is positioned between and
20 distracts apart adjacent spinous processes, allowing the spacer to come between the spinous processes. The spacer and guide can be moved in a posterior to anterior direction, and the spacer is able to rotate and translate in order to accommodate such movement. After this is accomplished, the first and second hook members are positioned through the incision and around upper and lower spinous processes.
25 Once that is accomplished, the spinous processes are urged towards each other and about the spacer, if this is not already the condition caused by the insertion of the spacer in order to distract the spinous processes. Then the racks of the shafts are meshed together, and the cap is placed upon the hub in order to secure the hooks firmly to the hub

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and thus to secure the spinous processes rigidly in position about the spinous processes.

Embodiments of Figs. 144-146c

5 A further embodiment 2300 of the invention is depicted in Fig. 144. In Fig. 144, the hub 2302 of this embodiment is depicted. This hub could be used, for example, with the embodiment shown in Fig. 141 and similar components are similarly numbered. In this embodiment, the hub 2302 includes an integral shaft 2324 with a threaded end 2325
10 which can accept a guide such as guide 2226 of Fig. 141. Unlike the embodiment in Fig. 141, this hub 2203 does not have a cap. Instead hub 2302 includes an open bore 2320 which is shaped in order to receive shafts 2310 and 2314, which have mating notches or teeth. Bore 2320 has a portion 2321 which is circular and which receives the
15 mated shaft 2310, 2324.

 Once this is accomplished, a screw 2322 is received in the threaded bore 2323 in order to lockingly position the mated shafts 2310, 2324. As this embodiment has an open bore 2320 and no cap, mating of the shafts 2310, 2314 to the open bore 2320 of the hub 2302 can be
20 done quickly and efficiently.

 Fig. 145 shows a hub 2402 of an embodiment 2400. This hub 2402 is similar to hub 2302, with the open bore 2420 having a shape which is different from the shape of bore 2320. In this embodiment bore 2420 includes a flat 2421 and a circular portion 2423. The shafts 2410
25 and 2414 when mated together would register in this open bore 2420. In particular, shaft 2414 has a flat which mates to flat 2421 and the combined shafts 2410 and 2424 have a circular portion which would mate to the circular portion 2423 of the bore 2420. Otherwise, hub 2202 would function similarly to hub 2302.

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Figs. 146a, 146b, and 146c depict a hub arrangement 2502 of an embodiment 2500 of the invention. In this embodiment, hub 2502 has two components 2511 and 2513. Component 2511 includes an open bore 2520 which is specially shaped in order to register shafts 2510, 2514 of the first and second hook members. In this particular embodiment, shaft 2510 is semi-circular in cross-section while shaft 2514 is triangular-shaped in cross-section. The triangular shape of shaft 2514 mates with the corner 2525 of the open bore 2520. The term open bore refers to 2520 and also to bores 2320 and 2420 in Figs. 144 and 145, and means that not only are both ends of the bore open, but there is a longitudinal slot along the length of the bore which is open, allowing access to the bore from the side of the bore. Once the shafts 2510, 2514 are inserted as shown Fig. 146c, a screw 2522 can be tightened through a bore of the hub 2502, locking the shafts in place. Once this has occurred, the first portion 2511 of the hub 2502 can be mated into the second portion 2513 of the hub 2502. In this embodiment, the second portion of the hub 2513 includes a slot 2515 into which can be slid or snapped into the first portion 2511. The first portion 2511 includes tangs 2517 and 2519 which fit under lips 2521, 2523 respectively as the first portion 2511 of the hub 2502 is slid or alternatively snapped into engagement with the second portion 2513. Once this occurs, a locking cam 2527 is turned in order to cause a cam member to be urged against this portion 2511 of the hub in order to lock 2511 to the second portion 2513. Alternatively, it is to be understood that the act of sliding or snapping hub portion 2511 into hub portion 2513 can be sufficient to lock portion 2511 into portion 2513. This embodiment further includes spacer 2536 and cone shaped guide 2526.

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Other features, functions, dimensions, and so forth of this embodiment are similar to the other embodiments as, for example, the embodiment of Fig. 141.

For purposes of insertion, one insertion methodology can be to
5 insert the second hub portion 2513 with the guide 2526 into the position
between the spinous processes. After this is accomplished, the hook
members can be positioned about the spinous processes and locked
into the first hub portion 2511. Then the first hub portion 2511 could be
slid or snapped into engagement with the second hub portion 2513.
10 Following that, the cam 2527 can be turned in order to secure the first
hub portion 2511 to the second hub portion 2513.

Embodiments of Figs. 147a-149b

Figs. 147a and 147b depict another embodiment 2600 of the
15 invention. This embodiment 2600 includes a hub 2602 and a rack and
pinion arrangement. The rack and pinion arrangement includes first and
second pinions 2660 and 2662. These pinions engage shafts 2610 and
2614 respectively. In these embodiments, these shafts 2610 and 2614
have rounded ends to which the hook is secured as depicted in, for
20 example, Fig. 131. For simplicity, these hooks have been left off of Figs.
147a, 147b. The position of the shafts 2610 and 2614 can be adjusted
relative to the hub. Once the shafts 2610, 2614 are appropriately
positioned the pinions can be locked in position, locking the shafts in
position. Pinions can be locked in position by tightening down screws
25 such as screw 2664 against the pinion 2660. A similar screw, not
shown, would tighten down pinion 2662.

Another embodiment of the invention, embodiment 2700 is
depicted in Figs. 148a and 148b. In this embodiment a bevel gear
arrangement 2770 is contained in the hub 2702. Bevel gear

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arrangement 2770 includes a first bevel gear 2772 and a second bevel gear 2774. Bevel gear 2772 has a shaft 2776 extending therefrom with a slot 2778. Slot 2778 can receive a tool for turning the bevel gear 2772. Bevel gear 2774 is mated to a threaded shaft 2710 of the hook member 2704. In this particular embodiment, the hook is not shown as is the case for the embodiment of Figs. 147a and 147b. When the bevel gear 2772 is turned, it turns bevel gear 2774. The turning of bevel gear 2774 causes the threaded shaft 2714 to retreat into or extend out of the center of the other shaft 2710. With the hook members positioned around spinous processes, the bevel gear 2772 can be used to turn bevel gear 2774 in order to draw the hook member 2706 toward the hub 2702, tightening the hook members about the spinous processes.

In this embodiment 2700, a shaft 2724 extends therefrom in order to receive a spacer and a guide in the same manner that, for example, the embodiment of Fig. 144 receives a spacer and a guide.

Figs. 149a and 149b depict embodiment 2800 of the invention. Embodiment 2800 includes a hub 2802 which houses a turnbuckle arrangement 2880 which is actuated by a worm gear drive 2882. Turnbuckle 2880 receives the threaded shaft 2810 and 2814 of the hook members 2804, 2806 respectively. As with the past embodiments, the actual hooks of these hook members are not depicted in order to simplify the drawing. By turning the turnbuckle 2880, the threaded shafts 2810, 2814 are either drawn into or urged out of the turnbuckle. Thus, by turning the worm gear 2882 with a tool placed in the slot 2884, the turnbuckle turns, causing the hook members to extend out of or be urged into the hub 2802.

Extending from the hub is a shaft 2824 with a threaded end 2825. As with the other embodiments, such as the embodiment in Fig. 144, a

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spacer can be placed on the shaft 2824 and a guide can be placed on the threaded end 2825.

5 The preferred method of inserting this embodiment is to insert the embodiment as a whole, placing the guide and spacer between the spinous processes. The hooks would be initially rotated 90° from their final orientation. Once inserted adjacent to the spinous processes, the hooks would be rotated by 90° and the spacer and the hooks would be further urged into contact with the spinous processes. Once this has occurred, the turnbuckle would be turned in order to tighten the hooks
10 about the spinous processes.

Embodiment of Fig. 150

Another embodiment 2900 of the invention is depicted in Fig. 150. This embodiment is similar to several of the other embodiments and, in particular, to the embodiment shown in Fig. 130. Accordingly, similar
15 elements will have similar least significant numbers. By way of example, the hub is designated 2902. In this particular embodiment, the hub is comprised of two components, the first hub component 2911 and the second hub component 2913. This is somewhat similar to the hub components shown in Fig. 146b.
20

The two hook members are secured to the first hub component 2911 in much the same manner as the hook members of Fig. 130 are secured to the hub in Fig. 130.

25 The hub 2902 is divided into first hub component 2911 and second hub component 2913 in order to add flexibility in the positioning of the guide and spacer fitted to second hub component 2913 with respect to the first and second hook members 2904 and 2906 which are secured to the first hub component 2911. Thus, should the anatomy of the spine and in particular the spinous process require, the spacer 2936

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and the guide 2926 can be moved relative to the first and second hook members 2904 and 2906 by selectively positioning the second hub component 2913 relative to the first hub component 2911. This can be accomplished by aligning the bore 2980 over one of the plurality of
5 bores 2982 positioned through the first hub component 2911. After this is accomplished, a threaded screw 2984 can be inserted through smooth bore 2980 and engage one of the threaded bores 2982 in order to secure the second hub component 2913 to the first hub component 2911, thus positioning the sleeve or spacer 2936 in a desired location
10 relative to the first and second hook members.

Embodiment of Fig. 151

Yet another embodiment of the invention 3000 is depicted in Fig. 151. Embodiment 3000 is meant for a double level spinous process
15 fixation. That is to say that three spinous processes are engaged and rigidly fixed together. Such a situation would occur, for example, when there is a double level primary fusion. That is, three adjacent vertebral bodies are all fused together. In such a situation a double level supplemental spine fixation device 3000 would be used. This
20 embodiment 3000 could be designed using any of the other embodiments depicted heretofore. Embodiment 3000 is in this particular instance modeled after the embodiment 2000 shown in Fig. 141. Accordingly, the elements that are similar to Fig. 141 have similarly least significant digits. By way of example, the hubs of Fig. 151
25 are both designated 3002 in accordance with the designation of Fig. 141. Similarly, the hub caps, sleeves, hook members, spacers, and guides are similarly numbered. In this embodiment two hubs, two spacers, and two guides are required as the first guide 3026 and the spacer 3036 would be inserted between first and second spinous

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processes, while the second guide 3026 and spacer 3036 would be inserted between the second and third spinous processes. The hook members 2004 and 2006 would hook about the first spinous process and the third spinous process respectively.

5 A preferred method of insertion of the device relative to three spinous processes would be to insert the guides and spacer between the first and second, and then the second and third spinous processes in order to distract apart the first and second spinous processes and also to distract apart the second and third spinous processes. After this
10 is accomplished, the first hook member would be placed about the first spinous process and the second hook member would be placed about the third spinous process. The shafts of the hook members would be inserted in the respective hubs 3002. In this situation, the shafts are both up-facing racks or teeth as shown in Fig. 151. A linking shaft 3039
15 has downwardly facing racks or teeth. Thus the upwardly facing rack or teeth of the first hook member 2004 would be laid in the upper hub 3002 with the teeth facing up. The teeth of the member 3039 facing down would engage the rack or teeth of the first hook 2004. Once this is accomplished, the cap will be placed over the hub and the screw
20 inserted in order to rigidly secure the hook member and the shaft 3039 relative to the upper hub 3002. Then the shaft of the second hook 2006 would be positioned in the lower hub 3002. The rack of shaft 3039 would mesh and lock with the rack of the shaft of the second hook member 2006. Once this is accomplished, the cap 3203 would be
25 placed over the hub and the screw would be inserted through the cap into the hub in order to secure the shaft 3039 and the second member 2006 relative to the lower hub.

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Industrial Applicability

From the above, it can be seen that the present invention can be used to successfully provide for supplemental spine fixation as an adjunct to primary spine fixation. Also spinous fixation without vertebral body fusion could be accomplished if that is desired. The embodiments of the invention provides the correct amount of rigidity between spinous processes with a minimally invasive device and methodology. The present invention does not require that structures associated with the spinous process, including bone and ligament, be altered for purposes of implantation, thus the device and method do not add to the trauma associated with spinal fusion.

Other features, aspects and objects of the invention can be obtained from a review of the figures and the claims.

It is to be understood that other embodiments of the invention can be developed and fall within the spirit and scope of the invention and claims.

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We claim:

1. An implant for rigidly positioning spinous processes comprising:
a first means adapted for engaging a first spinous process;
a second means adapted for engaging a second spinous process;
5 a body means adapted for positioning between the first spinous process and the second spinous process; and
a hub means for engaging the first means, the second means and the body means; and
wherein at least one of said hub means and said body means
10 includes means for allowing the body means to move relative to at least one of the first means and the second means.
2. The implant of claim 1 wherein:
said first means is a first hook;
15 said second means is a second hook; and
said body means includes a locking mechanism for locking the position of said first hook and said second hook relative to said body means.
- 20 3. The implant of claim 1 wherein:
said body means includes a locking mechanism for locking the position of said first means and said second means relative to said body means.
- 25 4. The implant of claim 3 wherein:
said allowing means includes a slot into which said body means is positioned in order to allow said body means to move relative to at least one of the first means and the second means.

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5. The implant of claim 1 wherein:
said body means includes an axis and a spacer positioned about
said axis, which spacer is rotatable about said axis.
- 5 6. The implant of claim 1 wherein:
said body means includes a guide that is adapted to guide the
body means between the first and the second spinous processes.
7. The implant of claim 1 wherein:
10 said allowing means allows the body means to move in a plane
which is substantially perpendicular to the first and second means.
8. The implant of claim 5 wherein:
said allowing means is located in said spacer and includes a bore
15 into which is received said axis so that said spacer can rotate about said
axis.
9. The implant of claim 5 wherein:
said allowing means is located in said spacer and includes a slot
20 with a said axis received in said slot such that said spacer can both
rotate about said axis and also move side to side relative to said central
axis.
10. The implant of claim 5 wherein:
25 said allowing means is located in said spacer and includes a slot
with a said axis received in said slot such that said spacer can both
rotate about said axis and also move in a direction substantially
perpendicular to said axis.

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11. The implant of claim 1 wherein:
at least one of said first means and said second means is shaped
in order to accommodate one of the spinous processes.
- 5 12. The implant of claim 1 wherein:
at least one of said first means and said second means has a
surface that is curved in order to mate with at least one of said spinous
processes.
- 10 13. The implant of claim 1 wherein:
at least one of said first means and said second means has a
surface which is shaped in order to allow at least one of the spinous
processes to mate with at least part of said surface.
- 15 14. The implant of claim 1 wherein:
at least one of said first means and said second means has a
surface which is shaped in order to allow at least one of said spinous
processes to mate with at least part of said surface without presenting
an abrupt surface change to the spinous process.
- 20 15. The implant of claim 1 wherein:
at least one of said first means and said second means is mounted
relative to said hub means so that at least one of said first means and
said second means can move relative to said hub means in order to
25 accommodate at least one of the first and second spinous processes.
16. The implant of claim 1 wherein:
at least one of said first means and said second means is swivel
mounted.

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17. The implant of claim 1 wherein:
said body means is elliptically shaped.

5 18. The implant of claim 1 wherein:
said body means is has an elongate shape.

19. The implant of claim 1 wherein:
said body means has a smooth continuous shape which is
adapted to be in contact with at least one of the spinous processes.
10

20. The implant of claim 1 wherein:
said body means has a smooth continuous shape which is
adapted to be in contact with at least one of the spinous processes and
wherein said shape is elongated.
15

21. The implant of claim 1 wherein:
said first means and said second means each have interlocking
means in order to allow said first means to interlock with said second
means.
20

22. The implant of claim 20 wherein:
said hub means includes a locking means for locking said first
means and said second means to said hub with said first means
interlocked with said second means.
25

23. The implant of claim 1 wherein:
said first means and said second means have first and second
interlocking means respectively;
said hub means has a third interlocking means; and

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said third interlocking means locks said first interlocking means and said second interlocking means to said hub means.

24. The implant of claim 1 wherein:

5 said hub means is shaped in a manner so that it can accept at least one of said first means and said second means in a certain orientation.

25. The implant of claim 1 wherein:

10 said hub means is shaped in a manner so that it can accept said first means and said second means in a certain orientation.

26. The implant of claim 1 wherein:

15 said hub means includes a first hub portion and a second hub portion;

 wherein said first hub portion and said second hub portion each include mating means for being mated together; and

 wherein said first hub portion has a means for engaging said first and second means and said second hub portion has means for
20 engaging said body means.

27. The implant of claim 1 wherein:

 said hub means includes a rack and pinion means for positioning at least one of said first means and said second means.

25

28. The implant of claim 1 wherein:

 said hub means includes a gear means for positioning at least one of said first means and said second means.

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29. The implant of claim 28 wherein:
said gear means includes a bevel gear.
30. The implant of claim 28 wherein:
5 said gear means includes first and second gears oriented about
perpendicular to each other.
31. The implant of claim 26 wherein:
said first hub portion is selective positionable relative to said
10 second hub portion in a plurality of positions.
32. The implant of claim 1 including:
a second body means for positioning between an intermediate
spinous process and one of the first spinous process and the second
15 spinous process; and
a second hub means for engaging the second body means, one
of said first means and said second means, and said hub means.
33. An implant for rigidly positioning spinous processes comprising:
20 a first hook adapted to engage a first spinous process;
a second hook adapted to engage a second spinous process;
a body adapted to be positionable between the first spinous
process and the second spinous process;
a hub;
25 said first hook, said second hook, and said body mounted to said
hub; and
wherein said body is movable relative to at least one of said first
hook and said second hook.

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34. The implant of claim 33 wherein:
said hub includes an axis; and
said body includes a bore, with said axis received in said bore in
order to allow said body to move relative to said axis.
- 5
35. The implant of claim 34 wherein:
said bore is cylindrical in shape.
36. The implant of claim 33 wherein:
said hub includes an axis; and
said body includes a slot, with said axis received in said slot in
order to allow said body to move relative to said axis.
- 10
37. The implant of claim 36 wherein:
said slot is shaped in order to allow said body to both rotate and
move side to said relative to said axis.
- 15
38. The implant of claim 36 wherein:
said slot is shaped in order to allow said body to both rotate and
move in a direction that is substantially perpendicular to said axis.
- 20
39. The implant of claim 33 wherein:
said hub includes a hub body and an axis which is movable
mounted to said hub body; and
said body is mounted on said axis.
- 25
40. The implant of claim 39 wherein:
said axis is partially received in a slot in said hub body so that
said axis can move in said slot relative to said hub body; and

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said body includes a bore, with said axis received in said bore.

41. The implant of claim 39 wherein:

said axis is partially received in a slot in said hub body so that

5 said axis can move in said slot relative to said hub body; and

said body includes a slot, with said axis received in said slot.

42. The implant of claim 33 wherein:

said hub includes a lock that locks the position of said first hook

10 and said second hook relative to said hub.

43. The implant of claim 33 wherein:

said body includes a guide that is adapted to guide the body
between the first and second spinous processes.

15

44. The implant of claim 33 wherein:

at least one of said first hook and said second hook is shaped in
order to accommodate one of the spinous processes.

20 45. The implant of claim 33 wherein:

at least one of said first hook and said second hook has a surface
that is curved in order to mate with at least one of the spinous
processes.

25 46. The implant of claim 33 wherein:

at least one of said first hook and said second hook has a surface
which is shaped in order to allow at least one of said spinous processes
to mate with at least part of said surface without presenting an abrupt
surface change to the spinous process.

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47. The implant of claim 33 wherein:

at least one of said first hook and said second hook is mounted relative to said hub so that at least one of said first hook and said second hook can move relative to said hub in order to accommodate at least one of the first and second spinous processes.

5

48. The implant of claim 33 wherein:

at least one of said first hook and said second hook is swivel mounted.

10

49. The implant of claim 33 wherein:

said body is elliptically shaped.

50. The implant of claim 33 wherein:

said body is has an elongate shape.

15

51. The implant of claim 33 wherein:

said body has a smooth continuous shape which is adapted to be in contact with at least one of the spinous processes.

20

52. The implant of claim 33 wherein:

said body has a smooth continuous shape which is adapted to be in contact with at least one of the spinous processes and wherein said shape is elongated.

25

53. The implant of claim 33 wherein:

said first hook and said second hook each have interlock in order to allow said first hook to interlock with said second hook.

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54. The implant of claim 53 wherein:

said hub includes a lock that locks for said first hook and said second hook to said hub with said first hook interlocked with said second hook.

5

55. The implant of claim 33 wherein:

said first hook and said second hook have first and second interlocks respectively;

said hub has a third interlock; and

10 said third interlock can lock said first interlock and said second interlock to said hub.

56. The implant of claim 33 wherein:

15 said hub is shaped in a manner so that it can accept at least one of said first hook and said second hook in a certain orientation.

57. The implant of claim 33 wherein:

said hub is shaped in a manner so that it can accept said first hook and said second hook in a certain orientation.

20

58. The implant of claim 33 wherein:

said hub includes a first hub portion and a second hub portion;

wherein said first hub portion and said second hub portion can be mated together; and

25 wherein said first hub portion can engage said first and second hooks and said second hub portion can engage said body.

59. The implant of claim 33 wherein:

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said hub includes a rack and pinion that can position at least one of said first hook and said second hook.

60. The implant of claim 33 wherein:

5 said hub means includes a gear arrangement that can position at least one of said first hook and said second hook.

61. The implant of claim 60 wherein:

10 said gear arrangement includes a bevel gear.

62. The implant of claim 60 wherein:

 said gear arrangement includes first and second gears oriented about perpendicular to each other.

15 63. The implant of claim 58 wherein:

 said first hub portion is selective positionable relative to said second hub portion in a plurality of positions.

64. The implant of claim 33 including:

20 a second body that is adapted to be positioned between an intermediate spinous process and one of the first spinous process and the second spinous process; and

 a second hub that engages the second body, one of said first hook and said second hook, and said hub.

25

65. In an implant for rigidly positioning spinous processes as an adjunct to spine fusion, the improvement comprising:
 a spacer positioned between adjacent spinous processes.

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66. The implant of claim 65 wherein:
said spacer is movably mounted on an axis so that said spacer
is movable relative to said axis at least while said spacer is implanted
between the spinous processes.

5

67. In an implant for rigidly positioning spinous processes as an
adjunct to spine fusion, the improvement comprising:
a spacer positioned between adjacent spinous processes; and
a first hook that is adapted to engage a first spinous process, and
10 a second hook that is adapted to engage a second spinous process.

68. The implant of claim 67 including:
a hub that connects said spacer and said first and second hooks.

15 69. The implant of claim 68 including:
said hub allows at least one of said spacer, said first hook and
said second hook to be selectively positioned relative to the others of
said spacer, said first hook, and said second hook.

20 70. The implant of claim 66 wherein:
said spacer has a slot, and said axis is received in said slot.

71. The implant of claim 69 wherein:
said slot allows said spacer to move substantially perpendicular
25 to said axis.

72. The implant of claim 68 wherein:
at least one of said first hook and said second hook can swivel.

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73. The implant of claim 65 including:
a second spacer positionable between spinous processes.
74. A method for rigidly positioning a first spinous process relative to
a second spinous process including the steps in any desired
order of:
placing a first hook around a first spinous process;
placing a second hook around a second spinous process;
using a hub to interlock the first hook relative to the second hook;
placing a spacer between the first spinous process and the
second spinous process; and
connecting said spacer to said hub.
75. The method of claim 73 wherein said connecting step includes:
connecting the spacer to the hub in a manner so that said spacer
can move relative to at least one of said first hook and said second
hook.
76. The method of claim 73 wherein said using step includes:
positioning said first hook relative to said second hook prior
interlocking the first hook relative to the second hook.
77. The method for rigidly positioning a first spinous process
relative to a second spinous process, including the steps in any desired
order of:
placing a first hook around a first spinous process;
placing a second hook around a second spinous process;

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placing a spacer between the first spinous process and the second spinous process, which spacer is mounted to the first portion of a hub; and

5 using a second portion of a hub to interlock the first hook relative to the second hook and then interlocking the first hub portion to the second hub portion.

78. The method of claim 74, including the step of not altering the first spinous process and the second spinous process as the method
10 is being performed.

79. The implant of claim 1, wherein at least one of said first means and said second means is selectively positionable relative to said hub means.
15

80. The implant of claim 1, wherein said implant does not require the alteration of the spinous processes for implantation with respect to the spinous processes.

20 81. The implant of claim 33, wherein at least one of said first hooks and said second hook is selectively positionable relative to said hub.

25 82. The implant of claim 33, wherein said implant does not require the alteration of the spinous processes for implantation with respect to the spinous processes.

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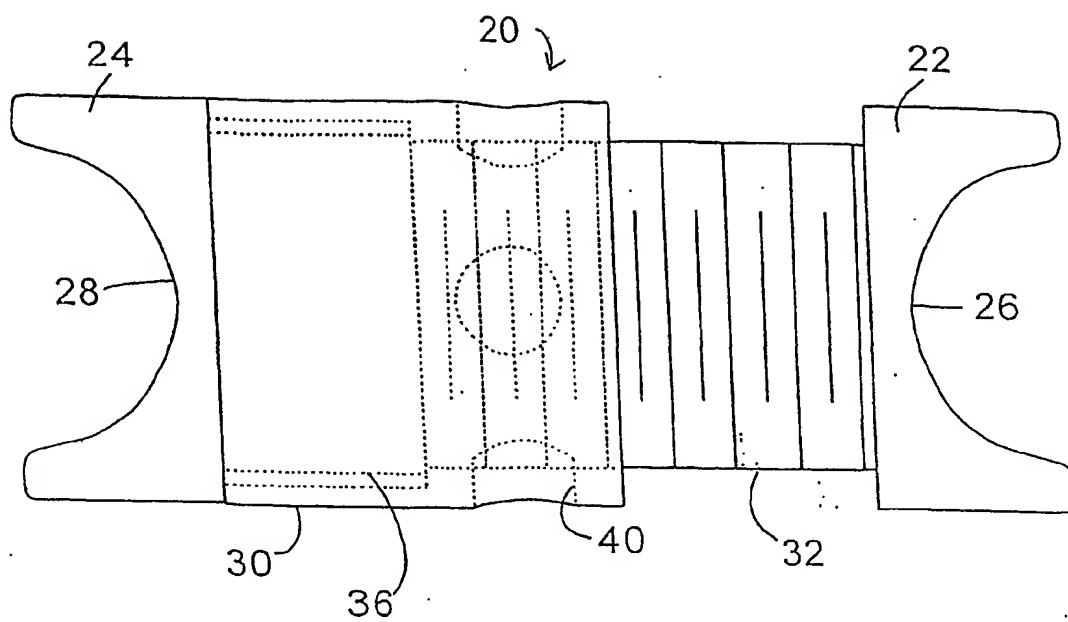


Fig. 1

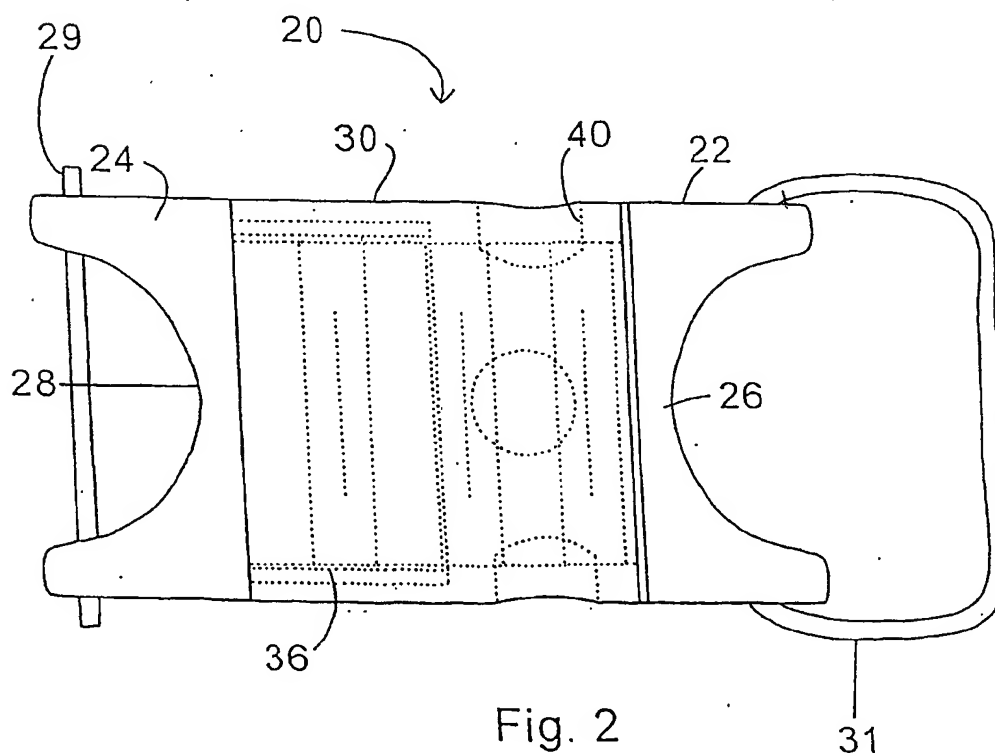


Fig. 2

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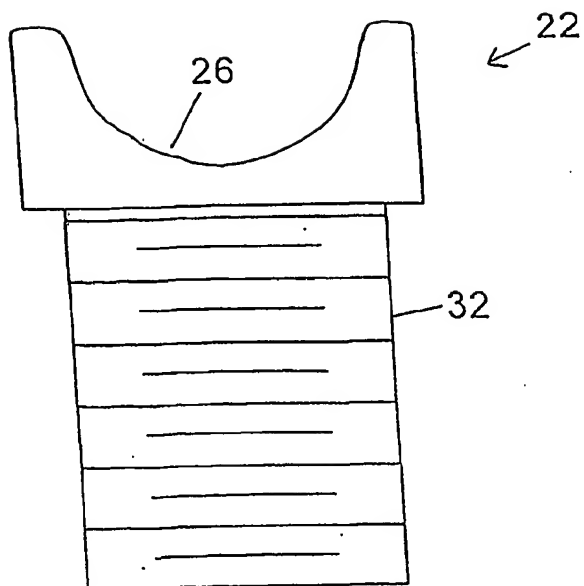


Fig. 3a

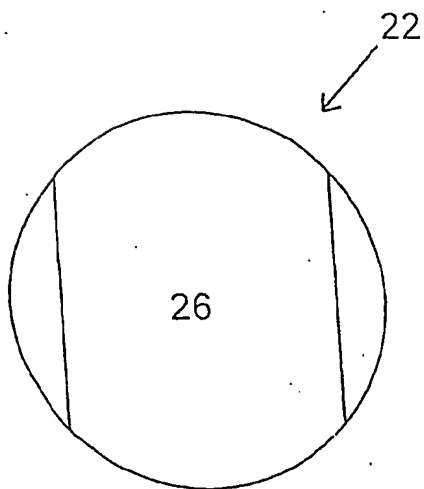


Fig. 3b

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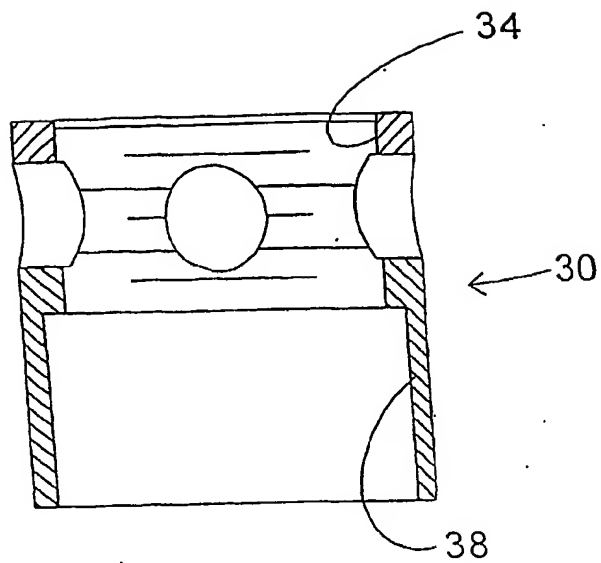


Fig. 4a.

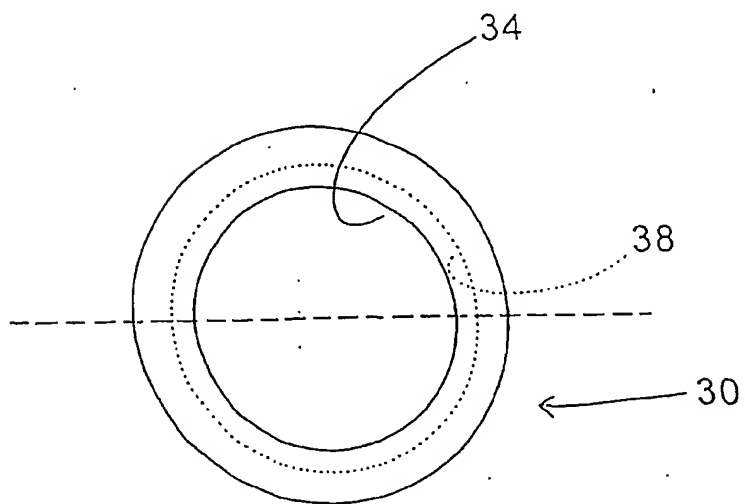


Fig. 4b

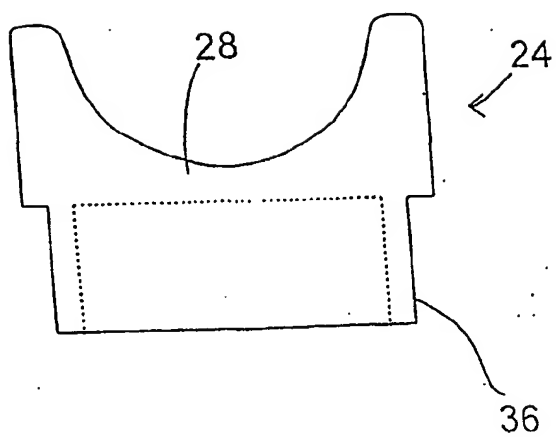


Fig. 5a

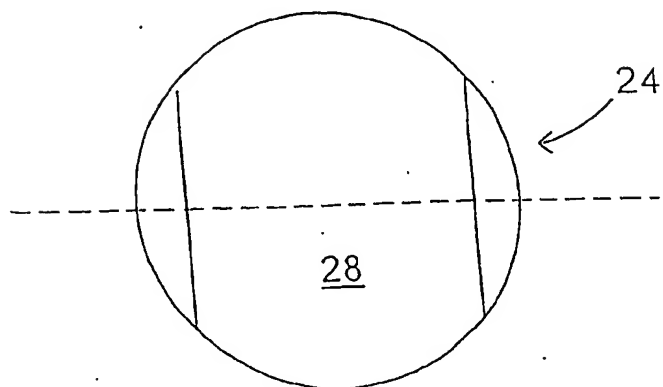
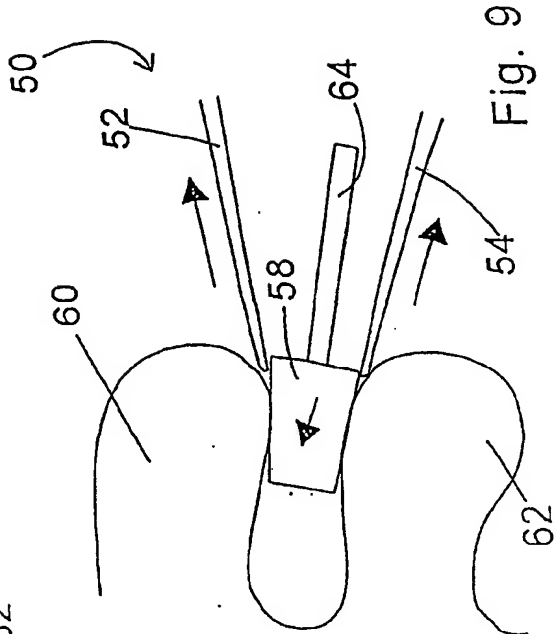
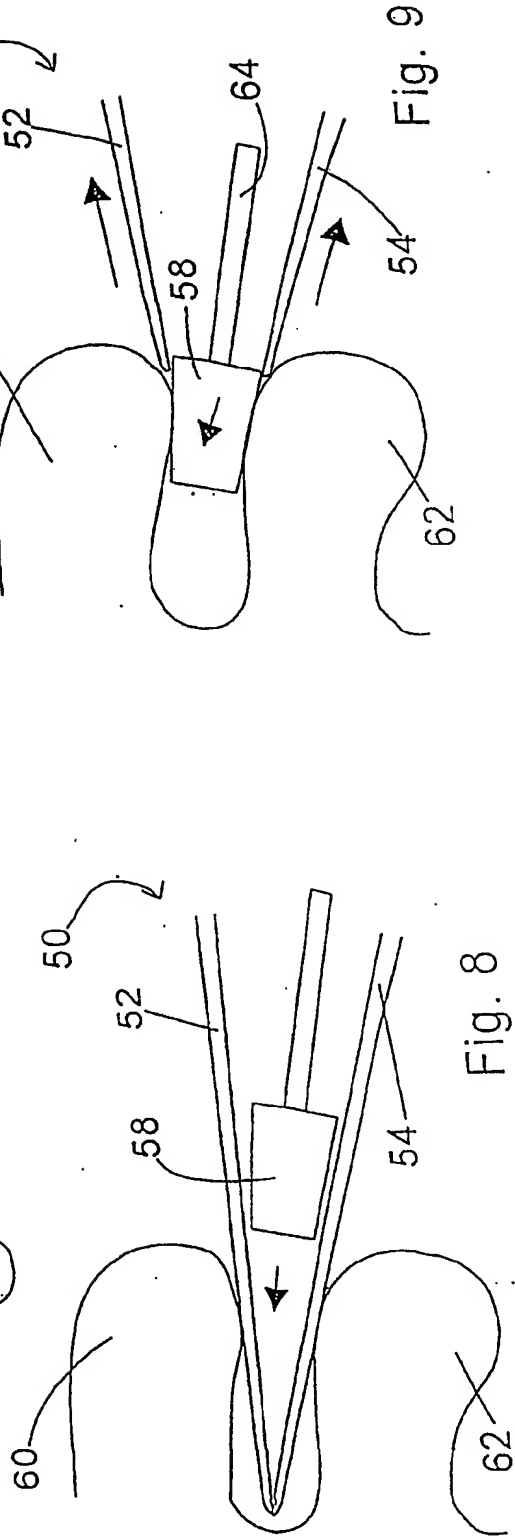
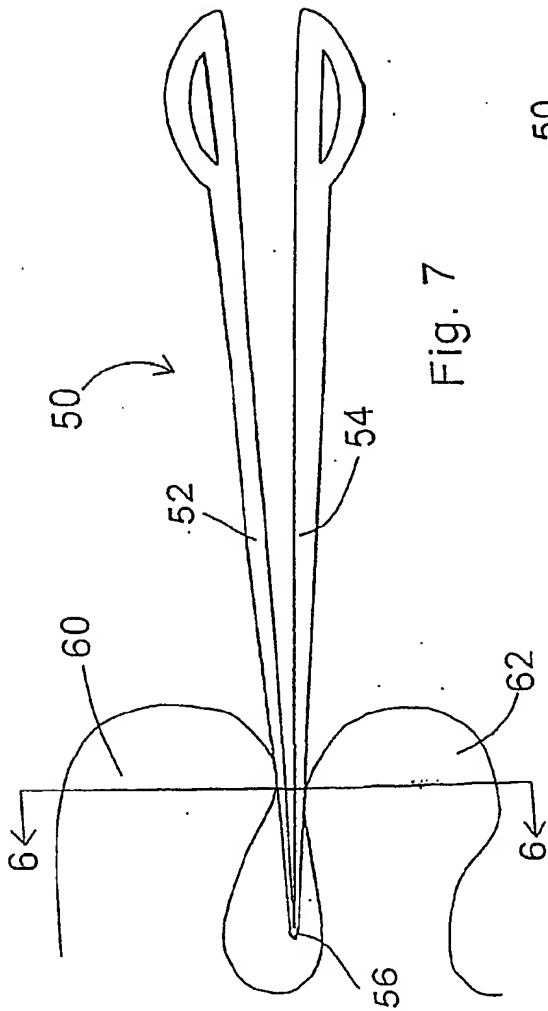
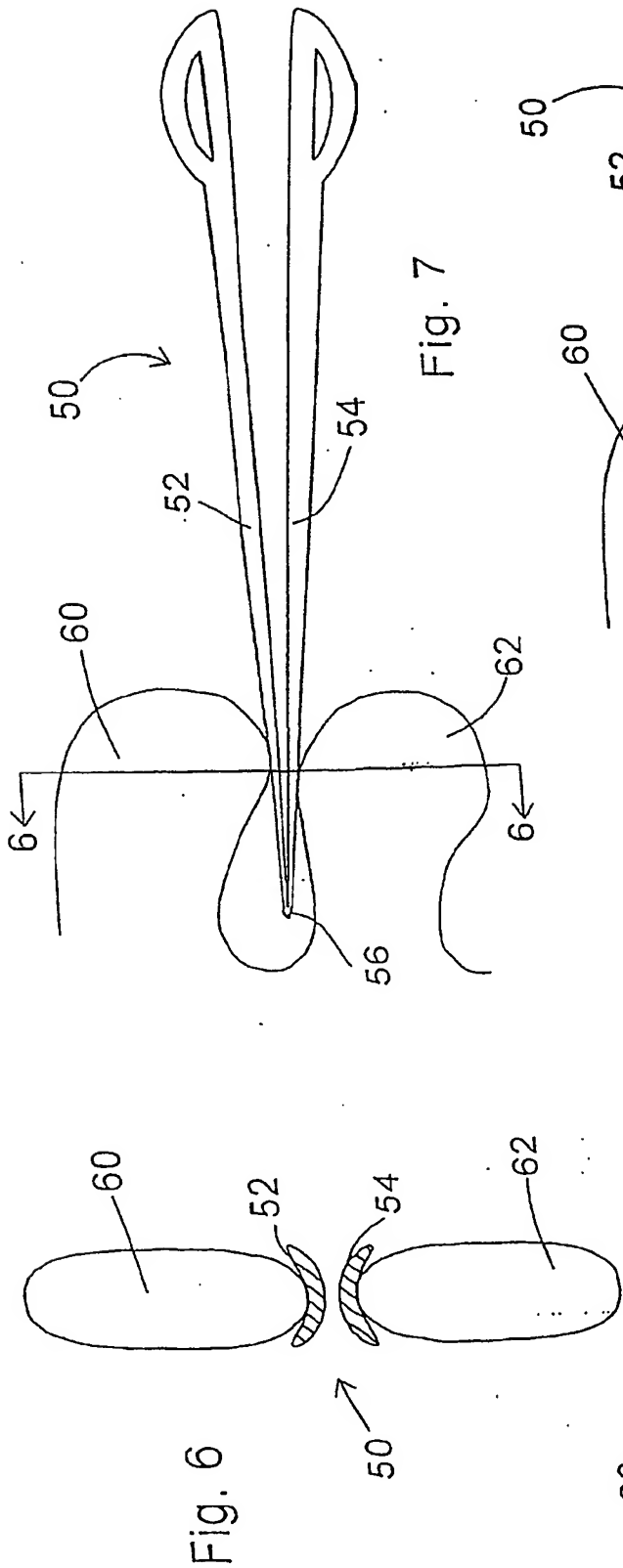


Fig. 5b



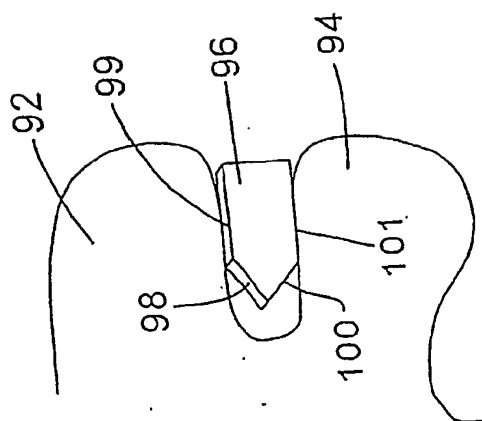


Fig. 11

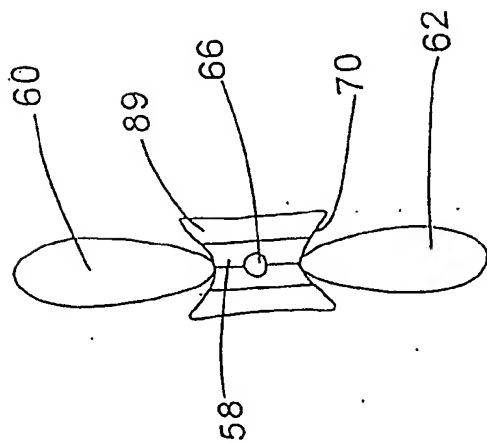


Fig. 10

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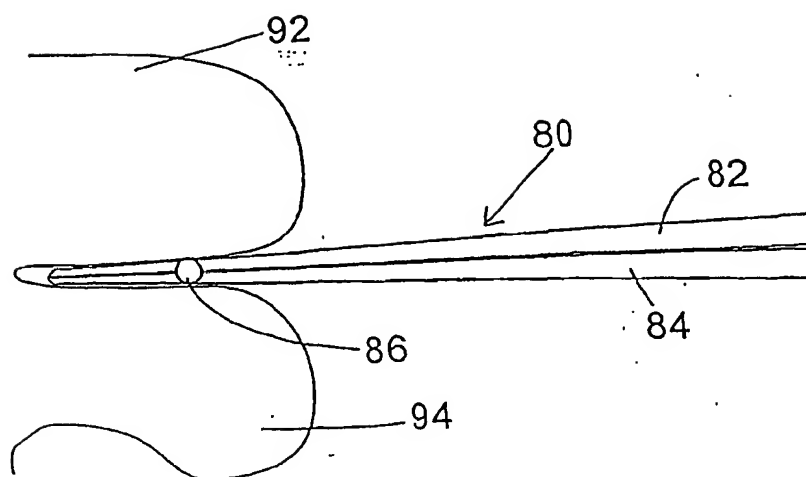


Fig. 12

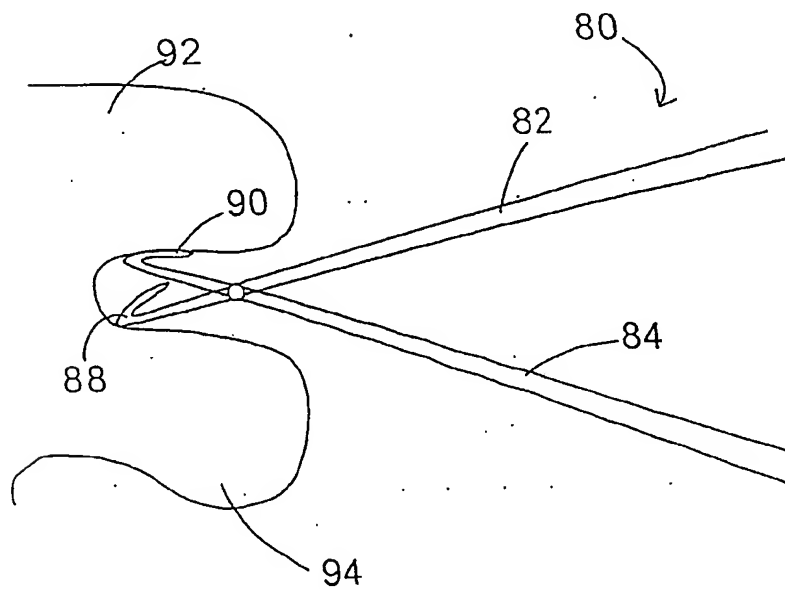


Fig. 13

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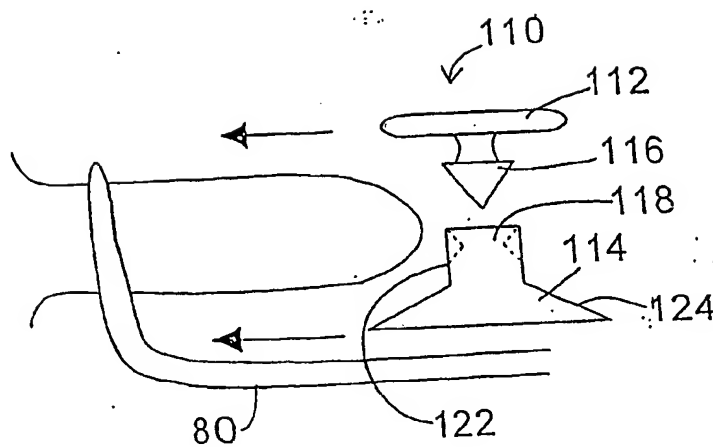


Fig. 14

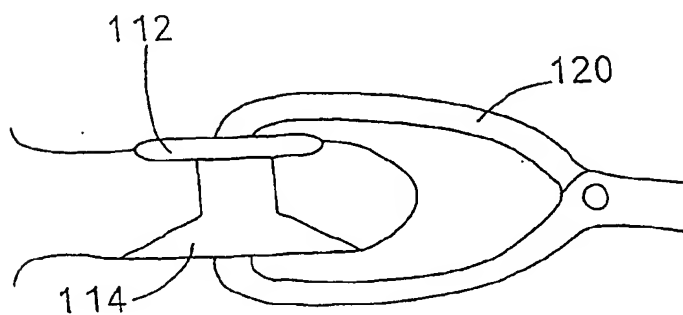


Fig. 15

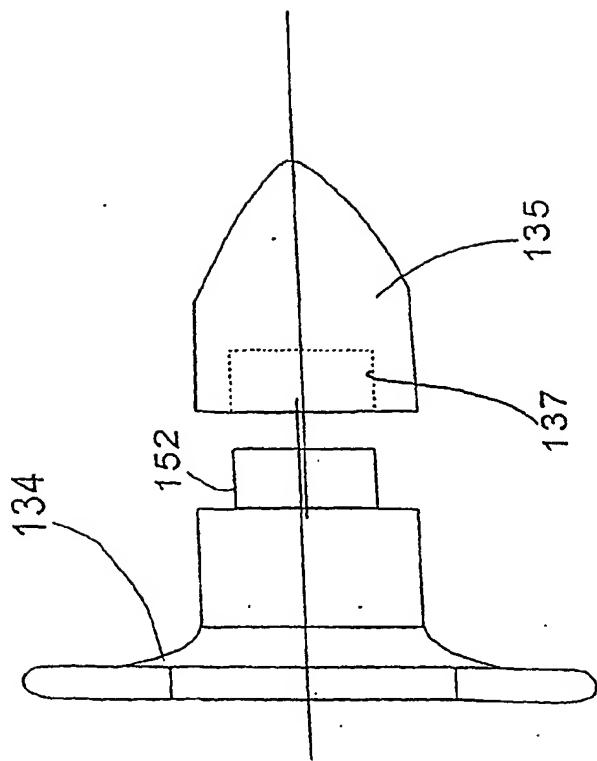
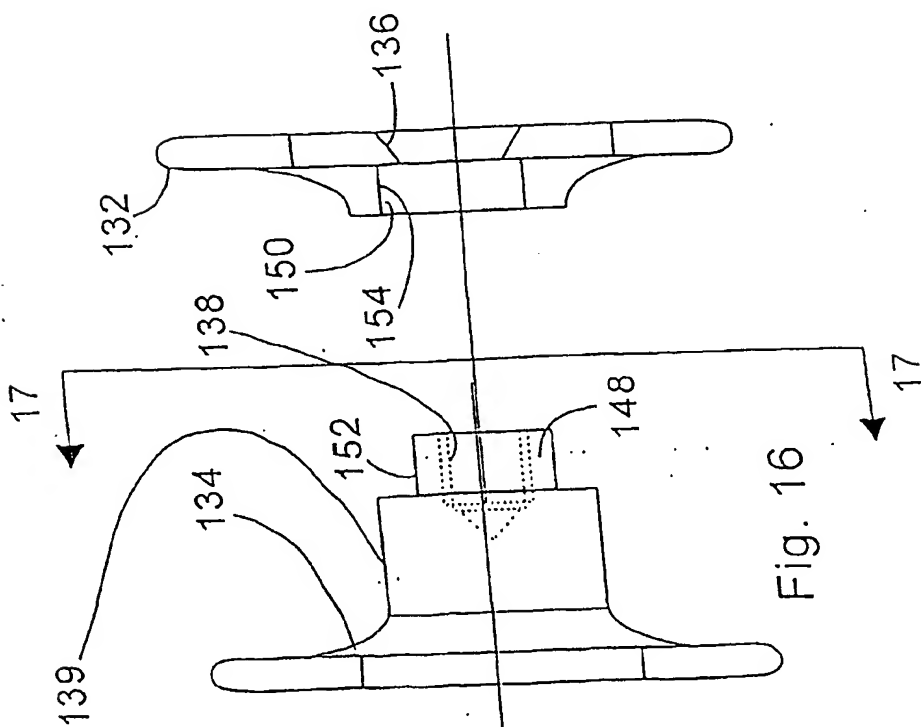
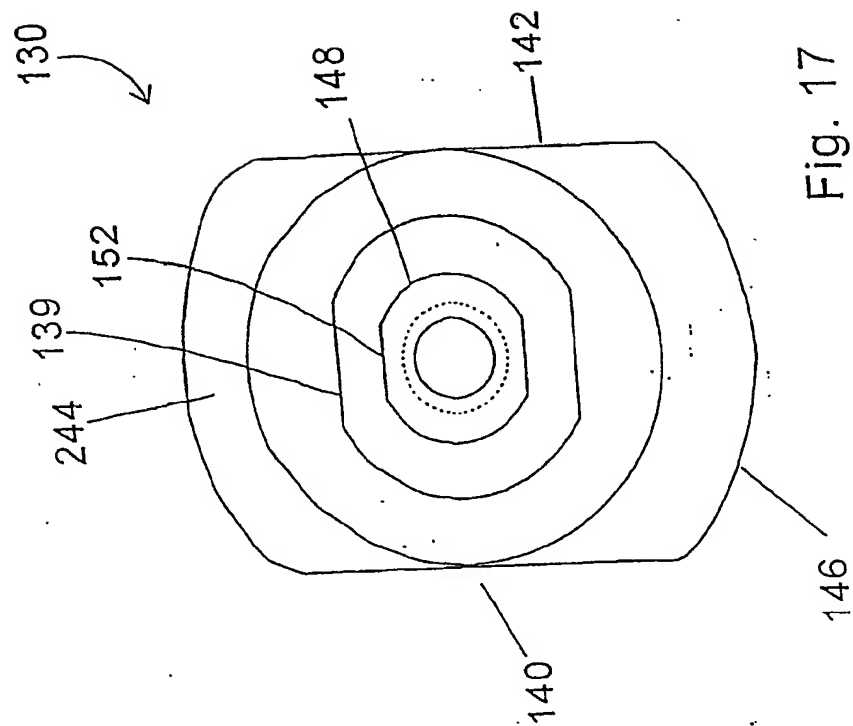


Fig. 16a

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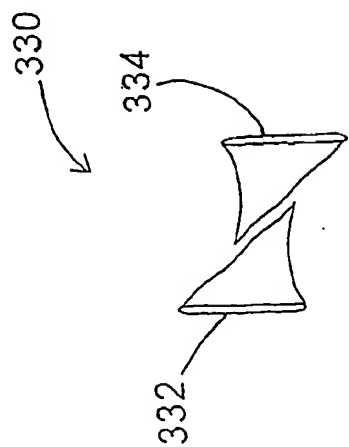


Fig. 18

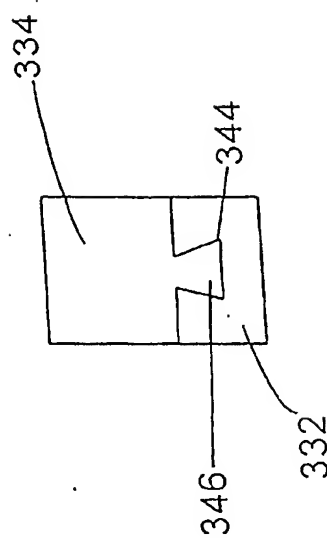


Fig. 19

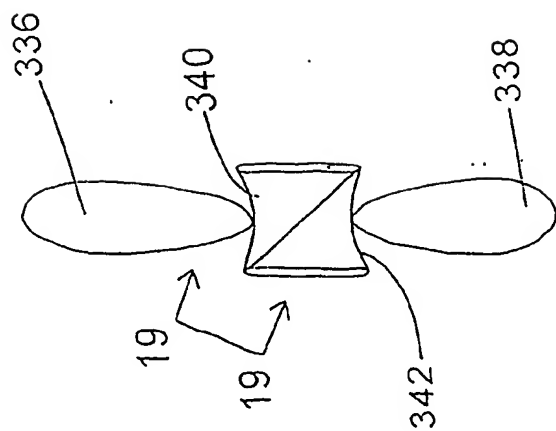


Fig. 20

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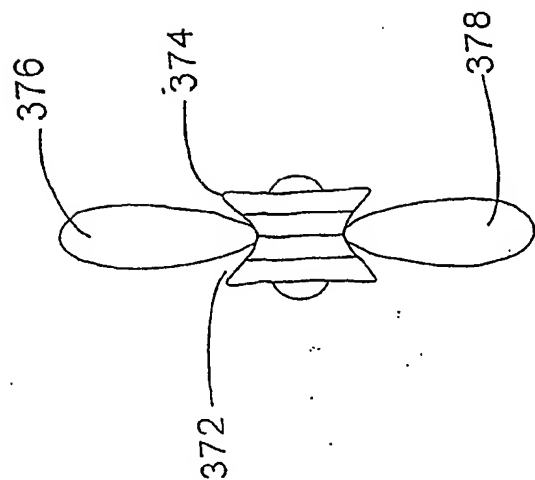


Fig. 22

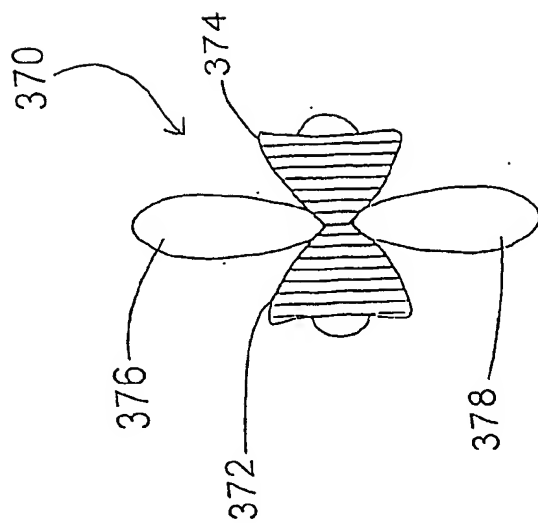


Fig. 21

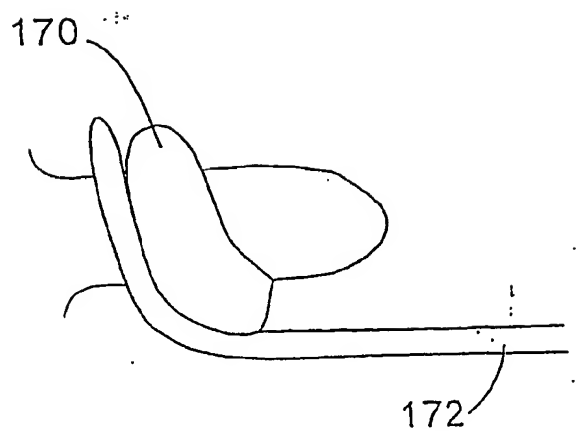


Fig. 24

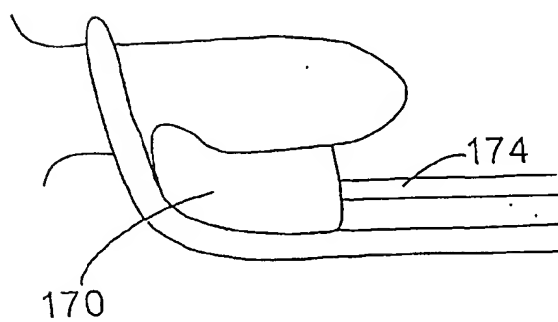


Fig. 23

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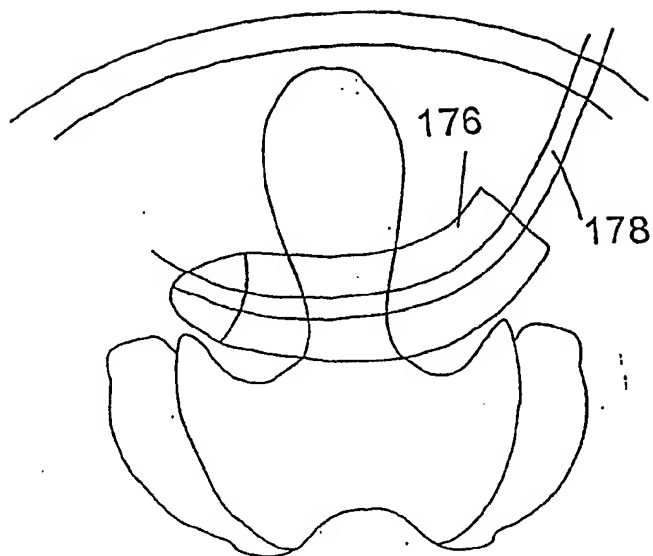


Fig. 25

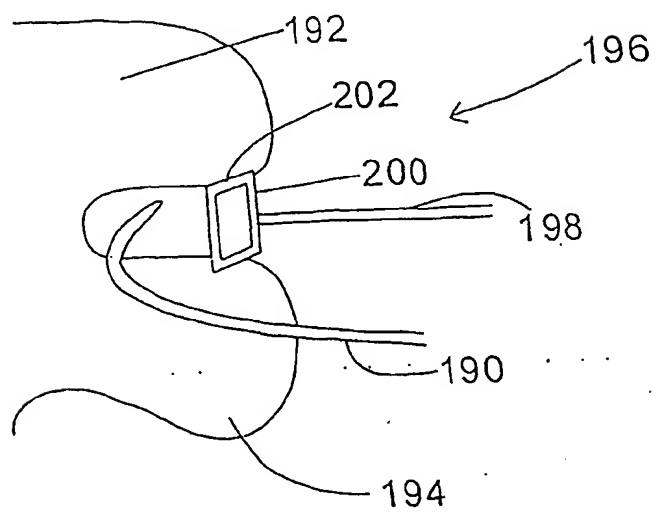


Fig. 26

Fig. 28

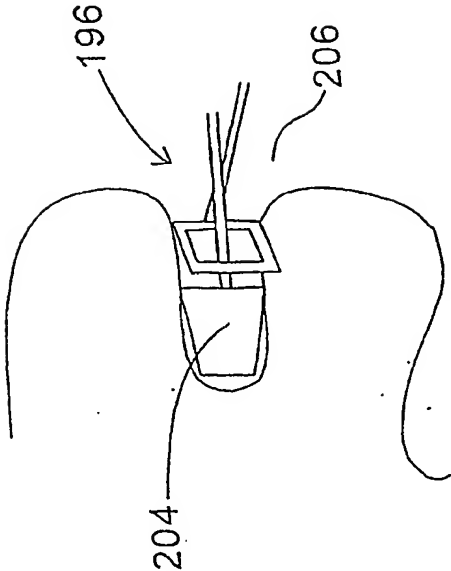


Fig. 30

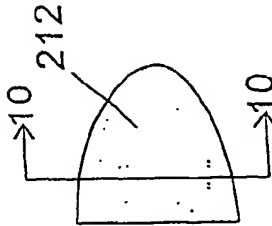


Fig. 27

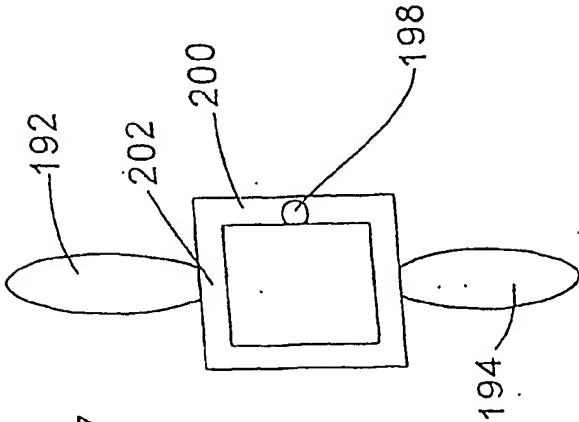
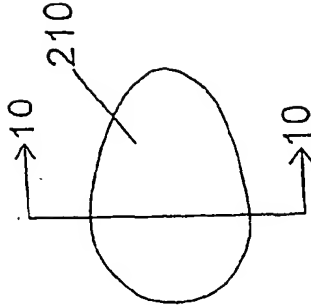


Fig. 29



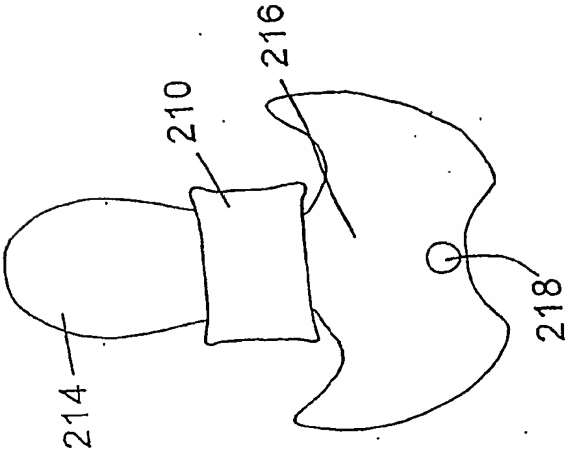


Fig. 33

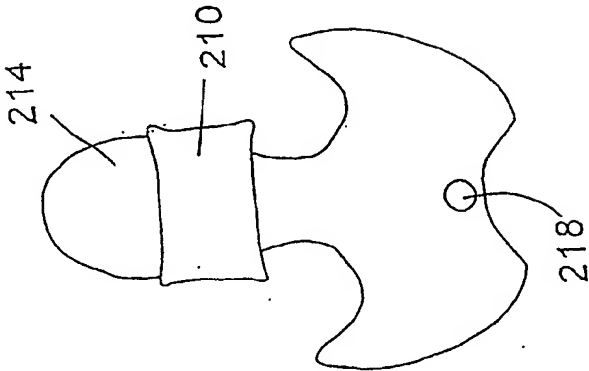


Fig. 32

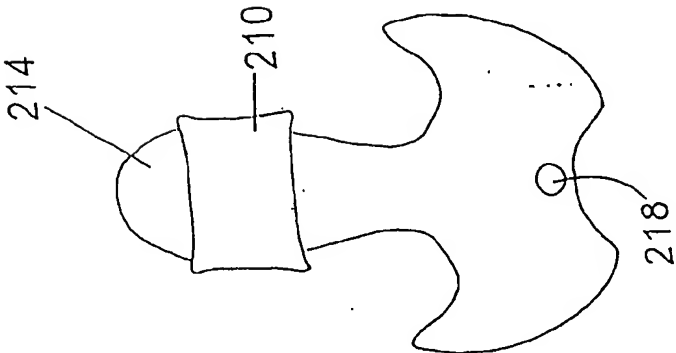


Fig. 31

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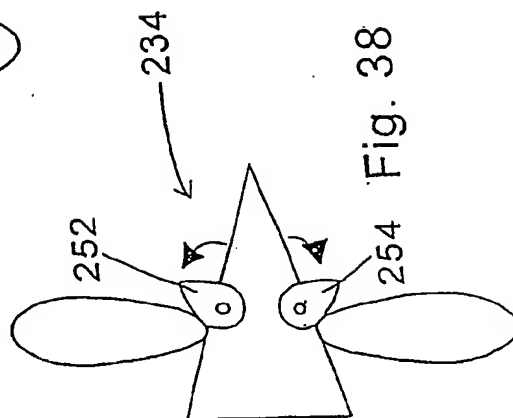
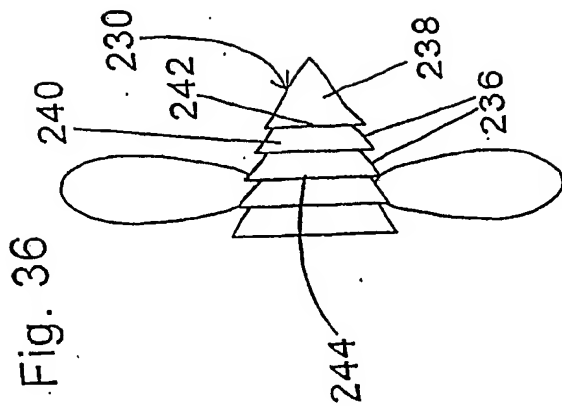


Fig. 35

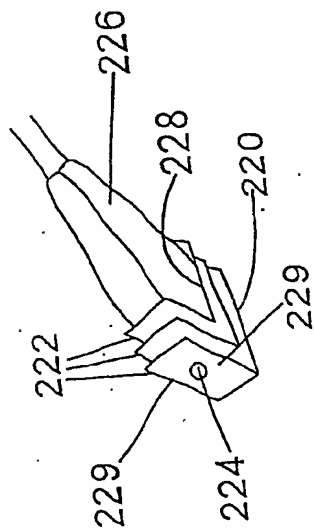


Fig. 34

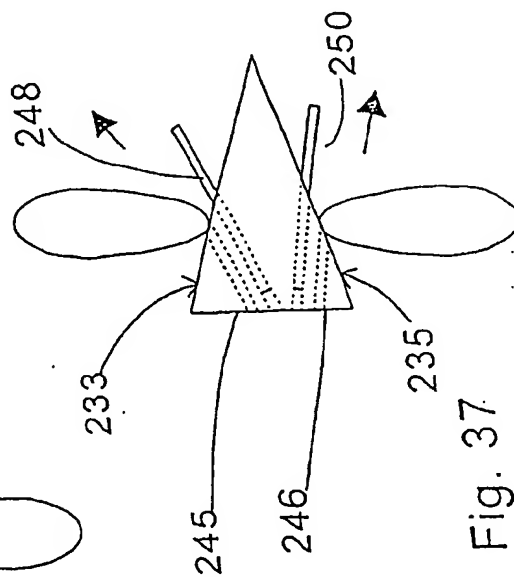
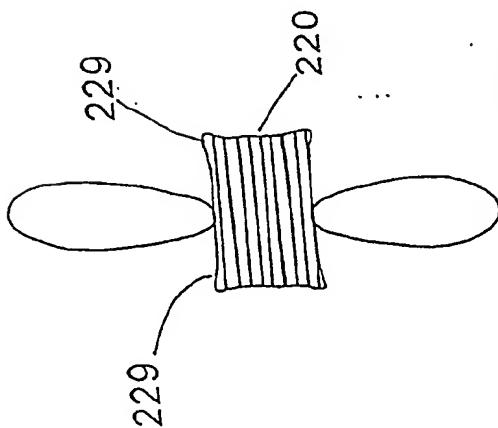
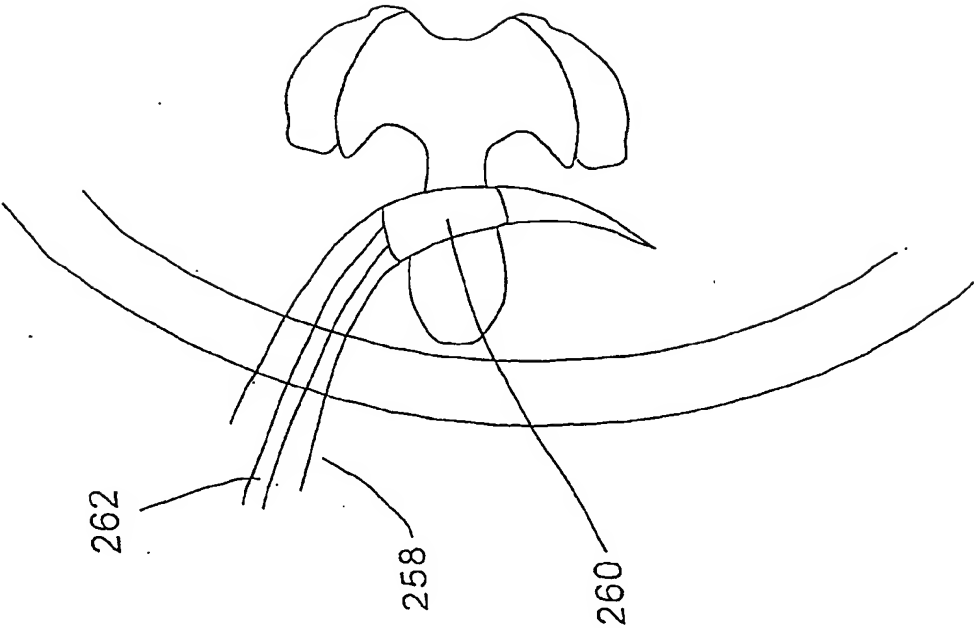


Fig. 37

Fig. 39



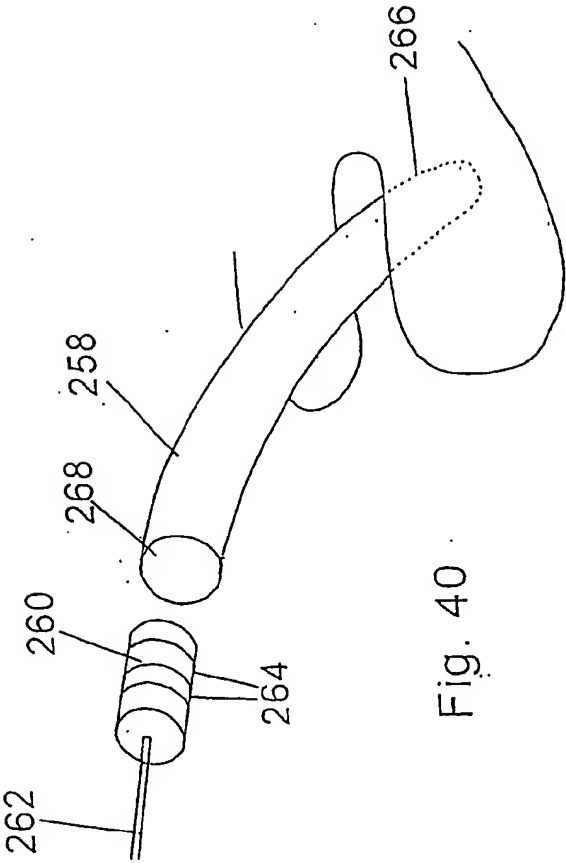


Fig. 40

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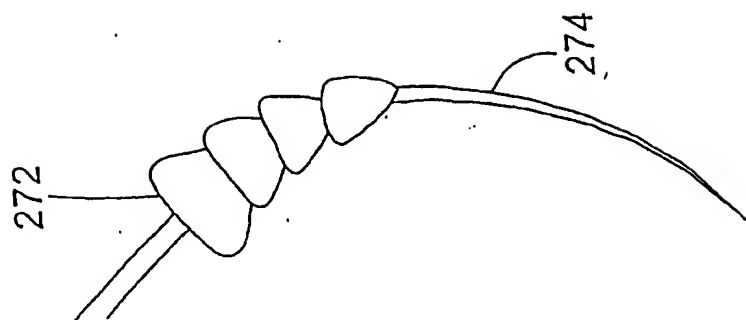


Fig. 43

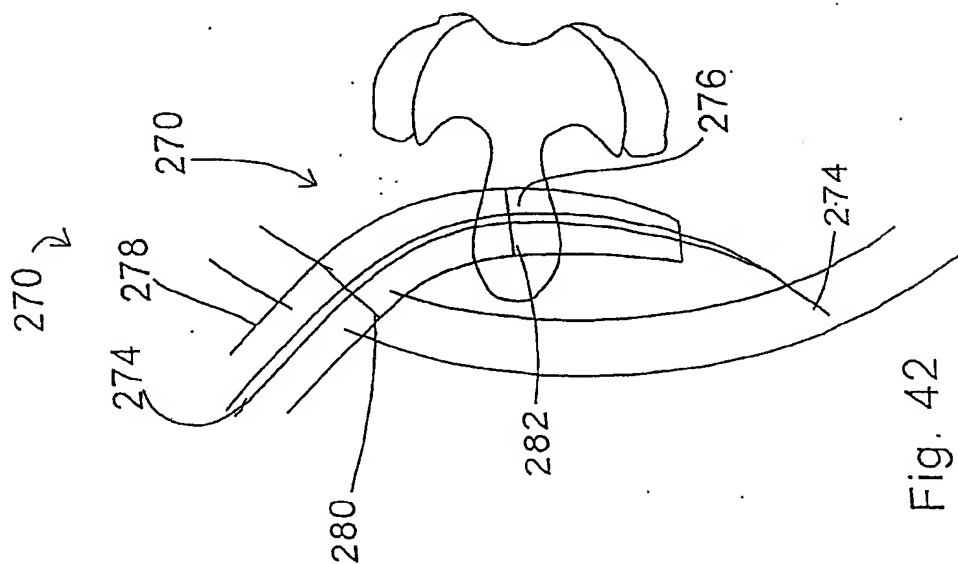


Fig. 42

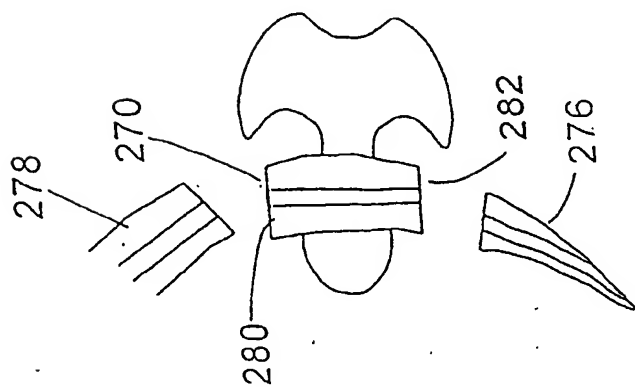


Fig. 41

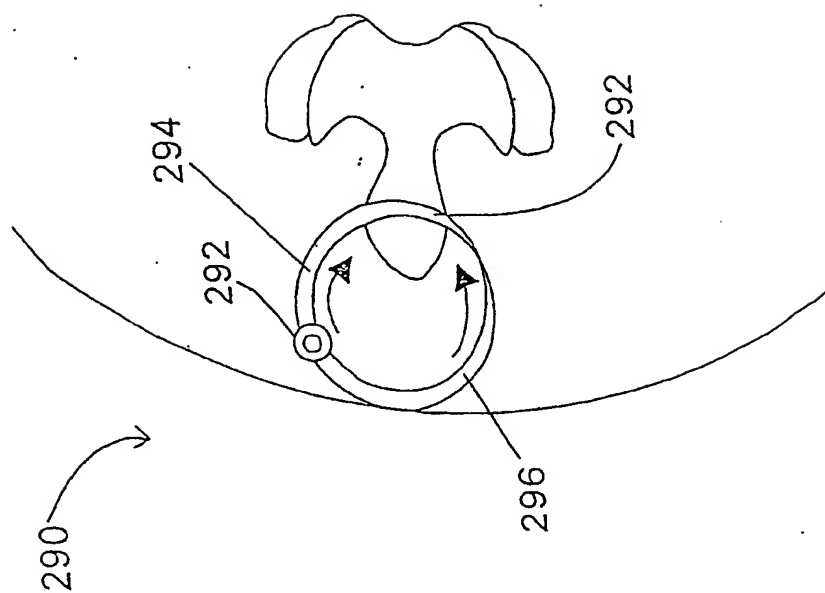
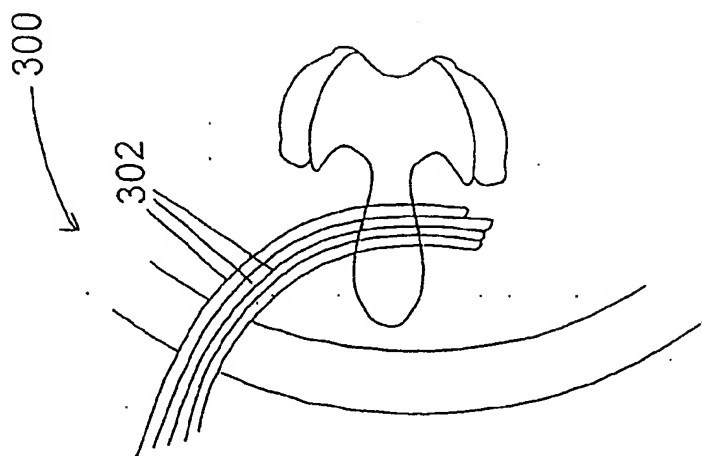
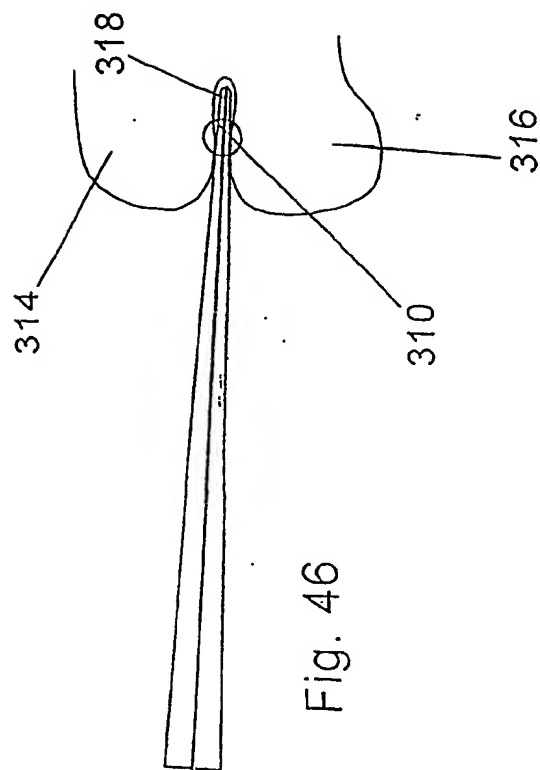
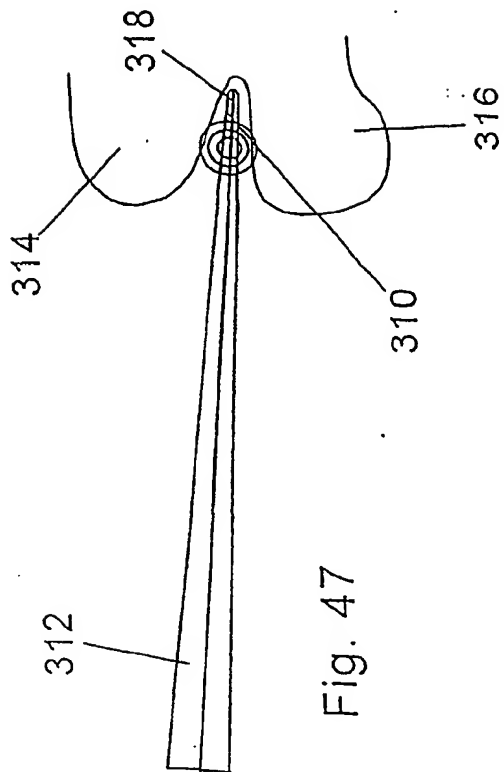


Fig. 44

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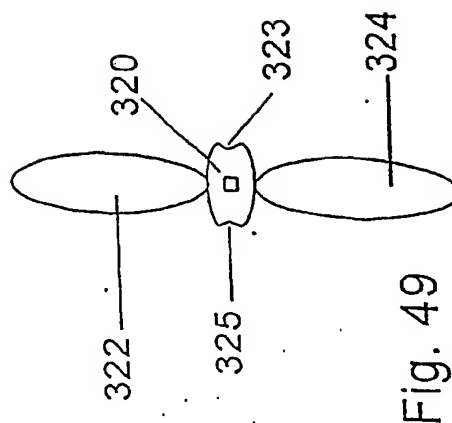
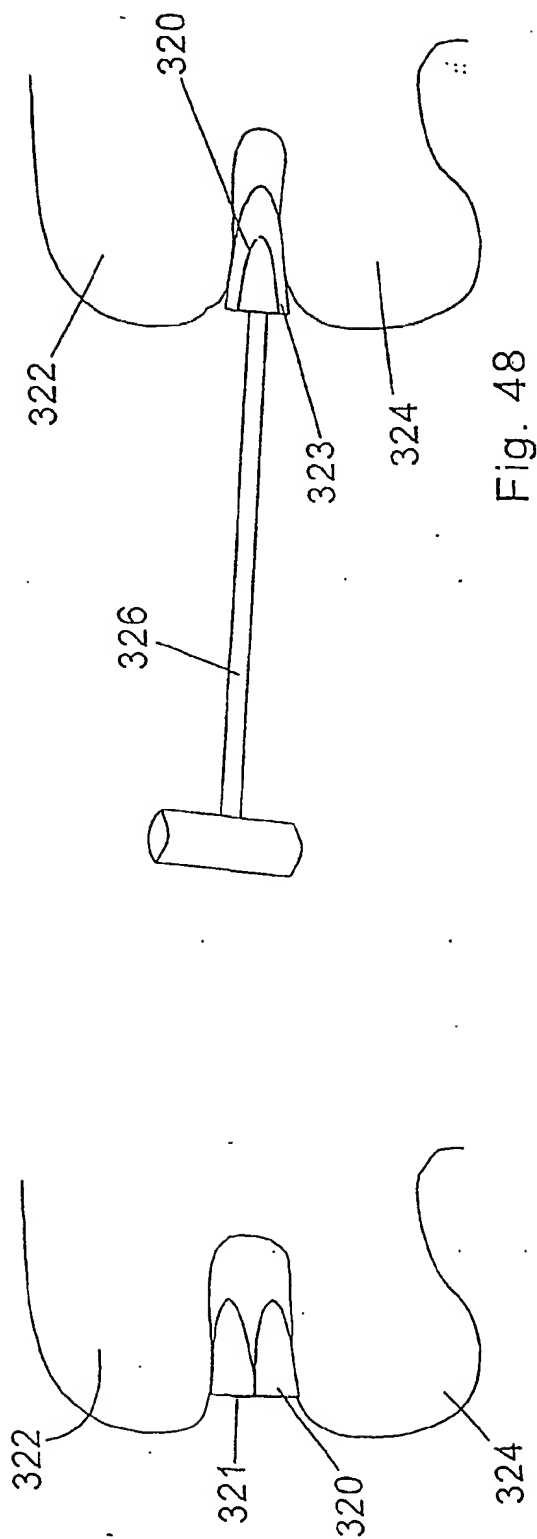


Fig. 51

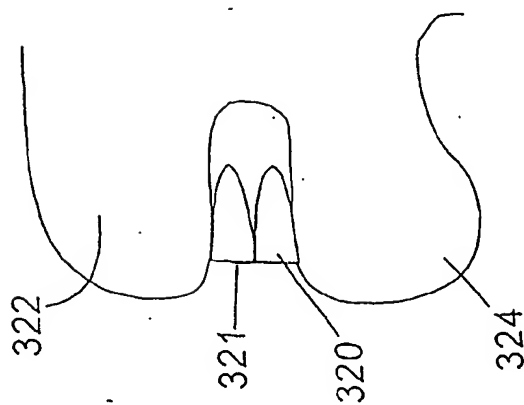
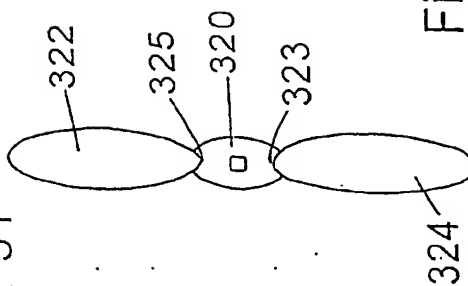


Fig. 50



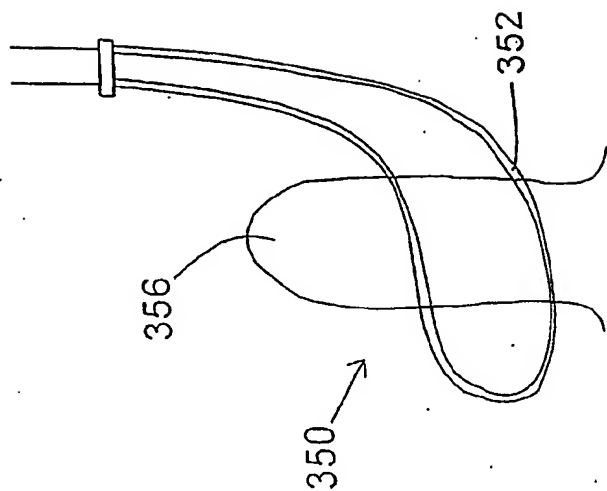


Fig. 54

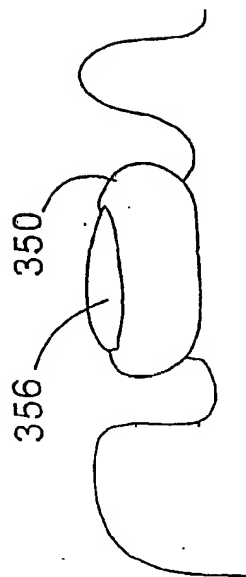


Fig. 55b

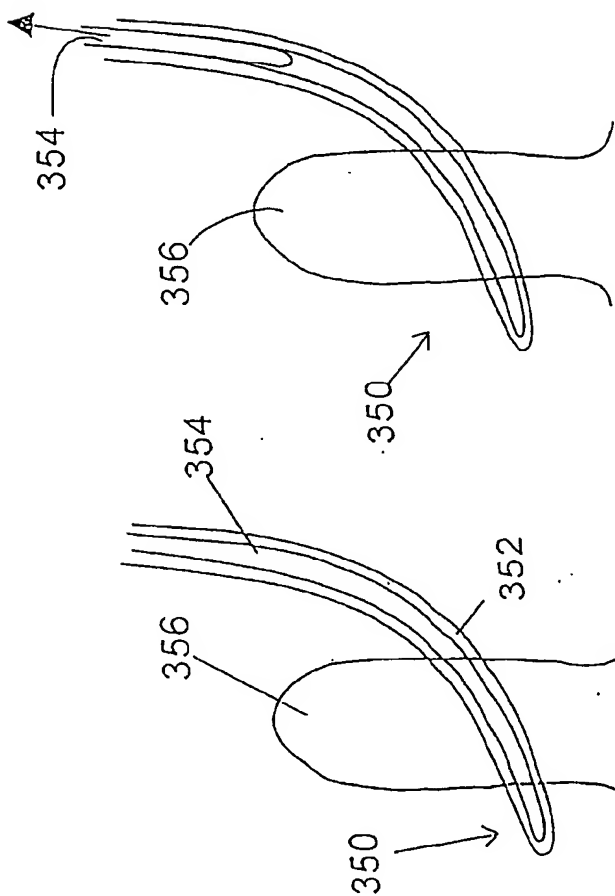


Fig. 53

Fig. 52

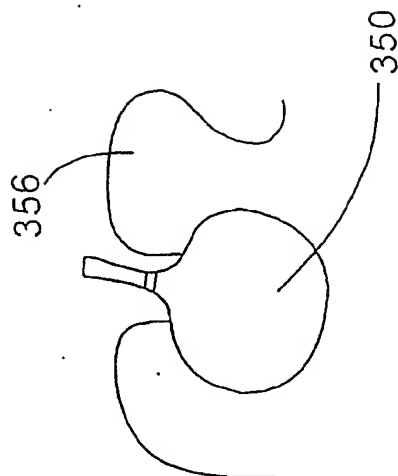


Fig. 55a

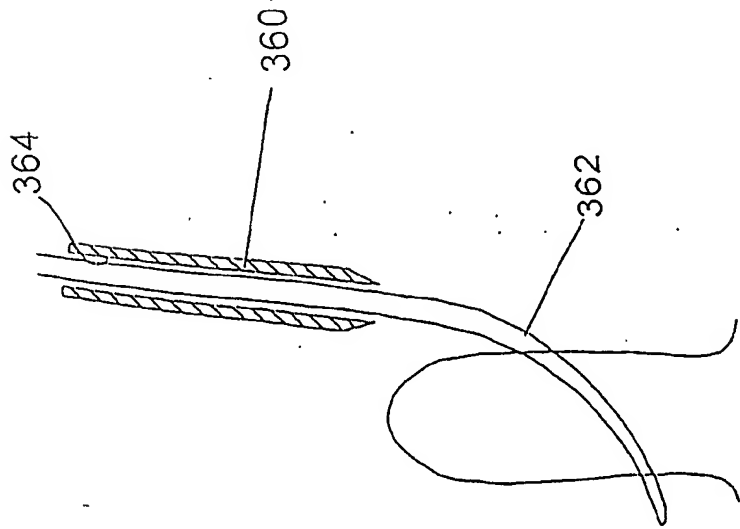


Fig. 56

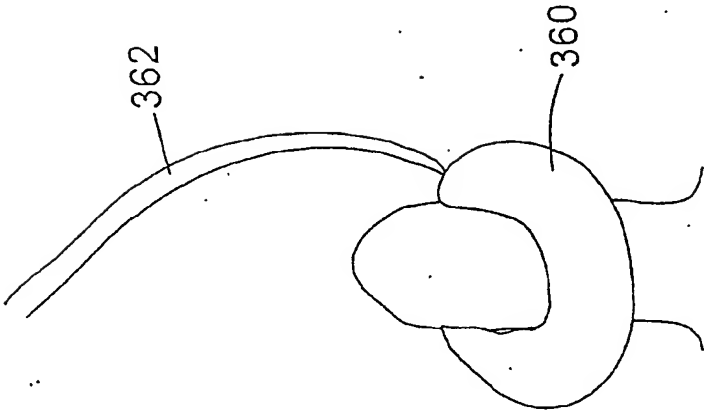


Fig. 57

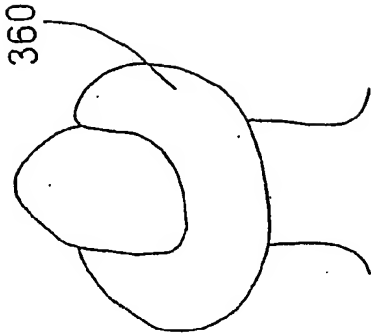


Fig. 58

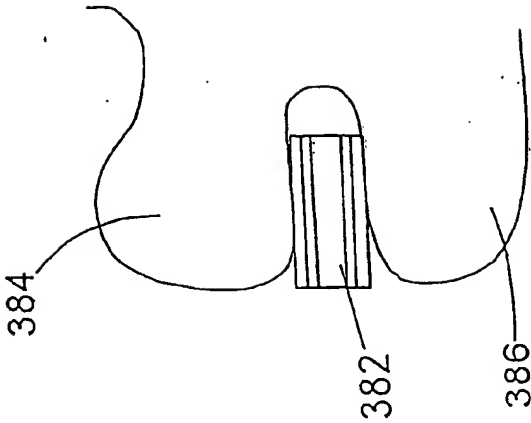


Fig. 60

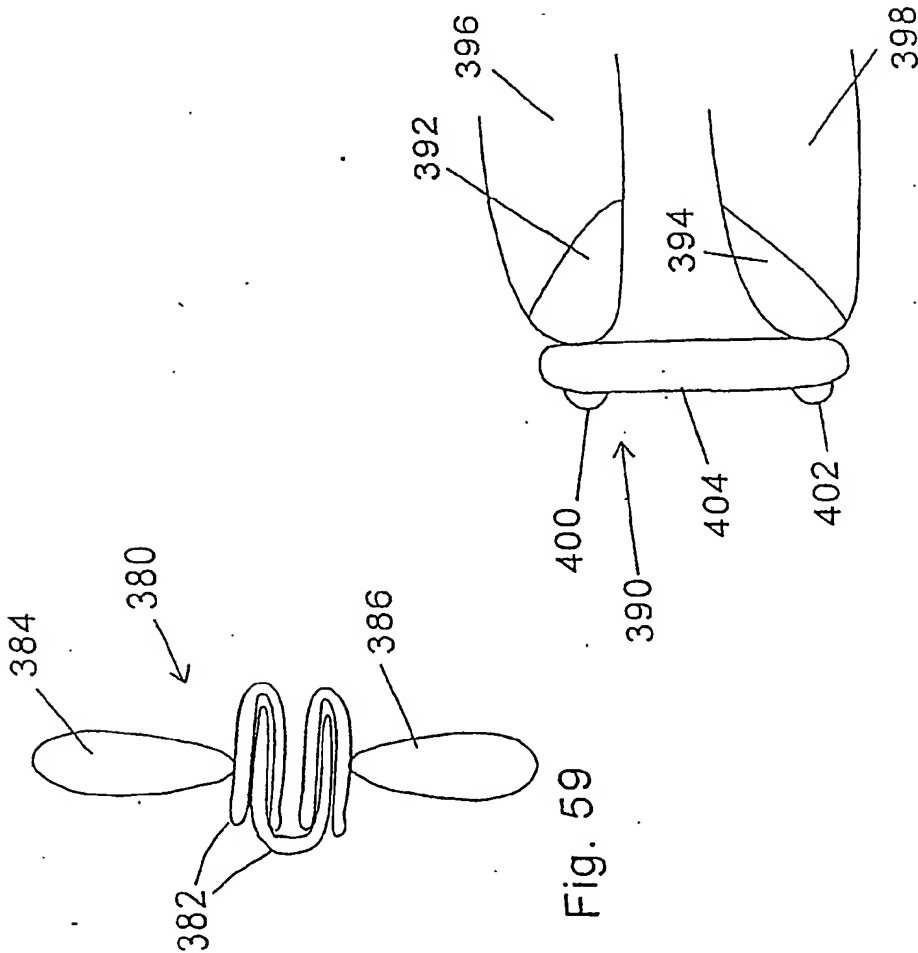


Fig. 59

Fig. 61

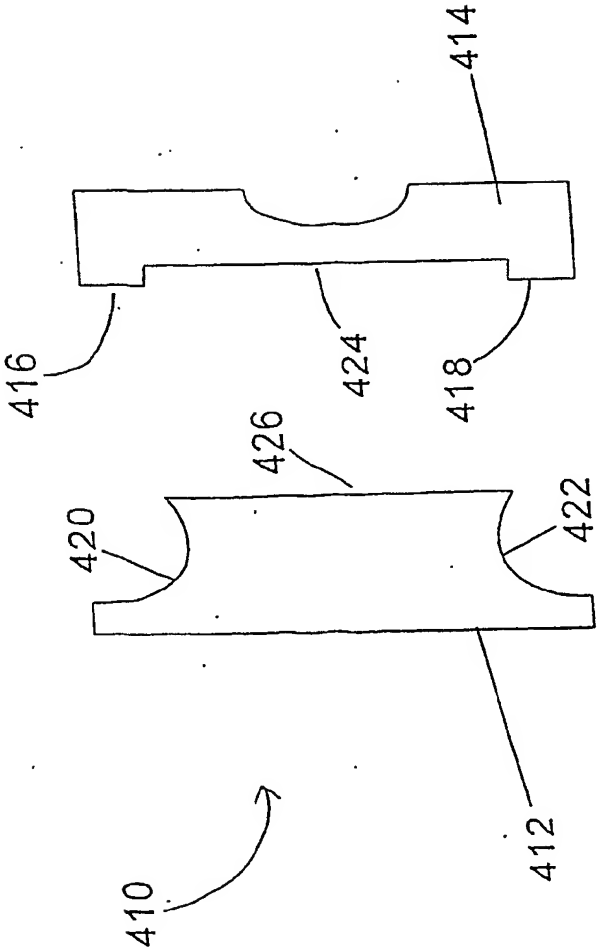


Fig. 62

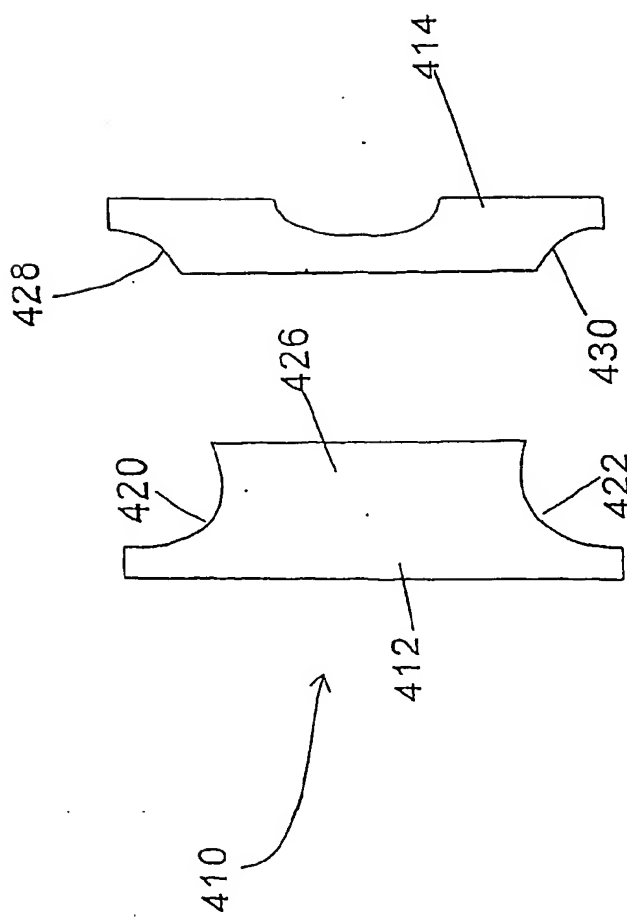


Fig. 63

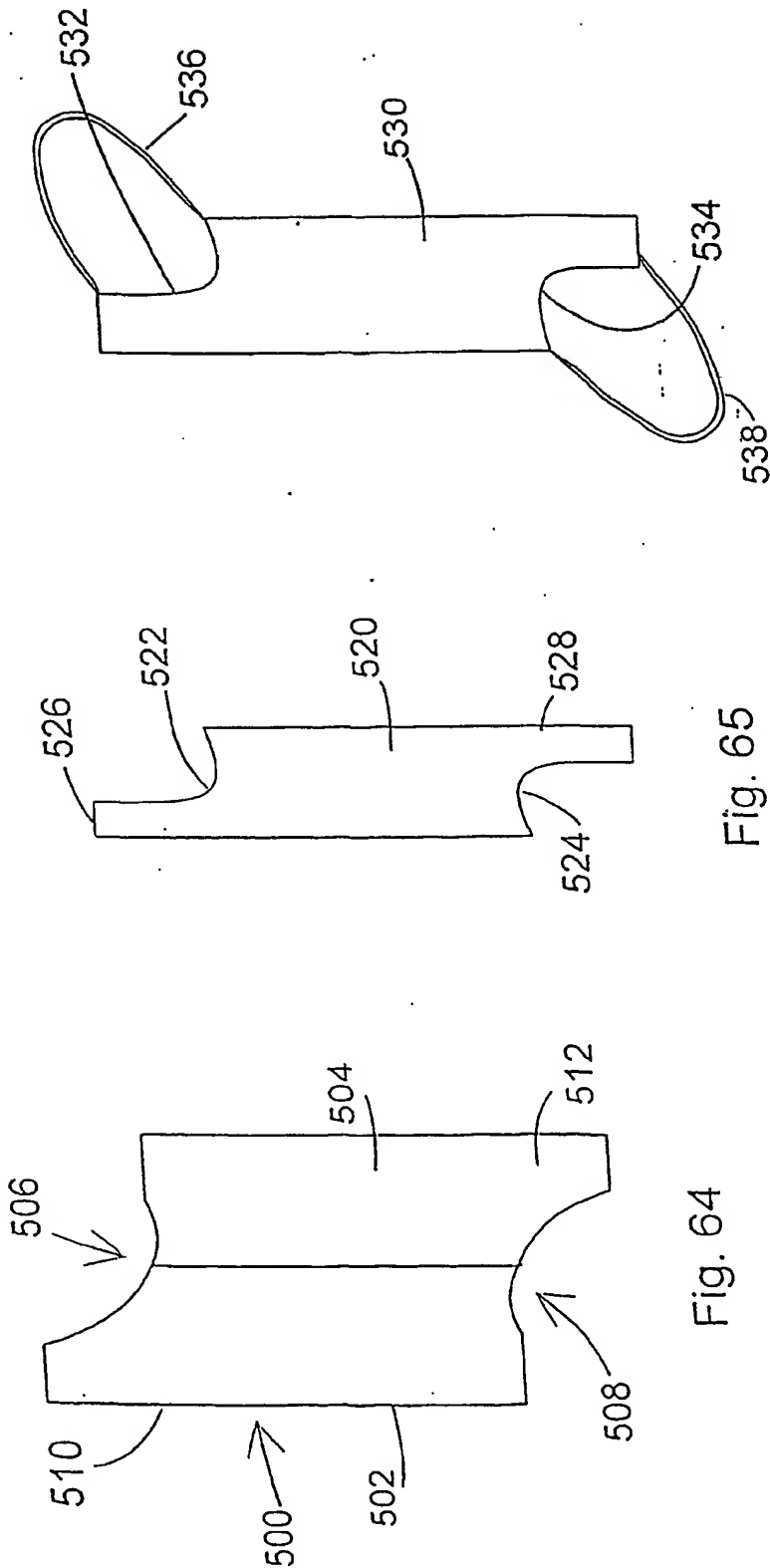


Fig. 64

Fig. 65

Fig. 66

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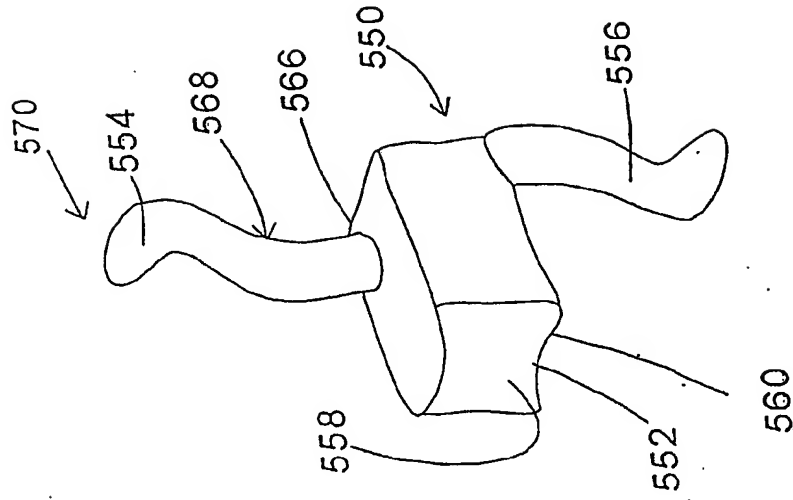


Fig. 68

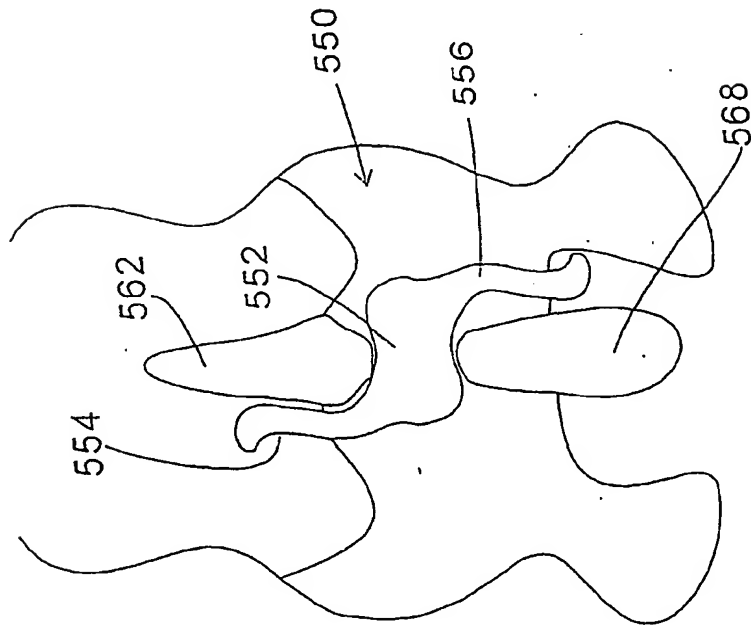


Fig. 67

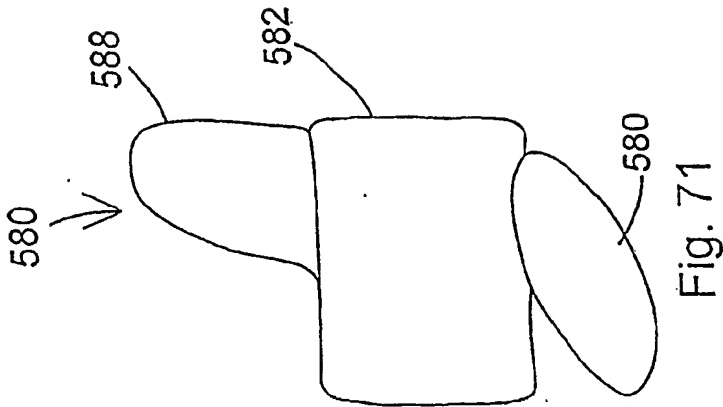


Fig. 71

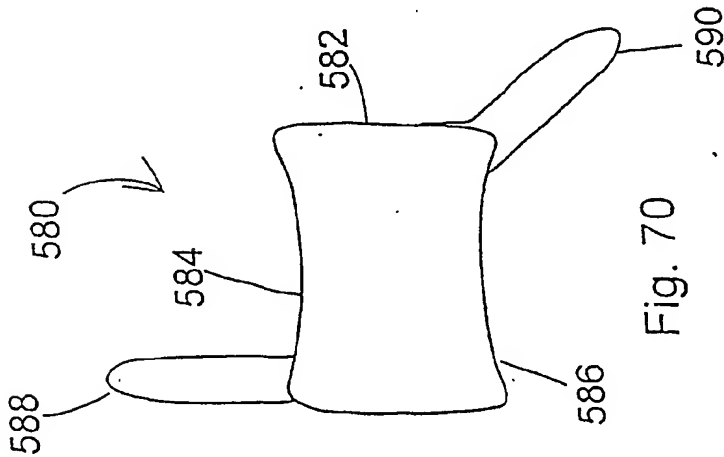


Fig. 70

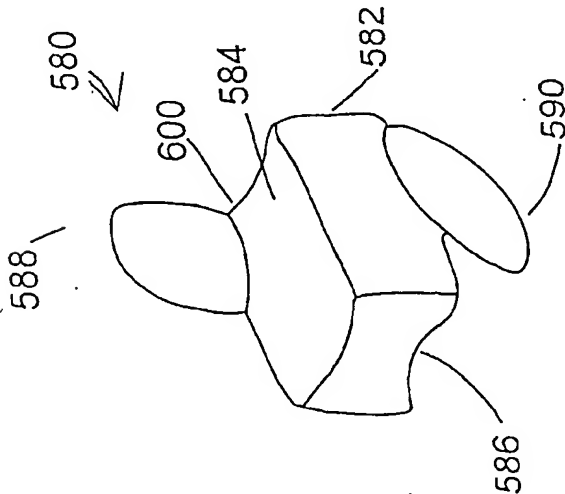


Fig. 69

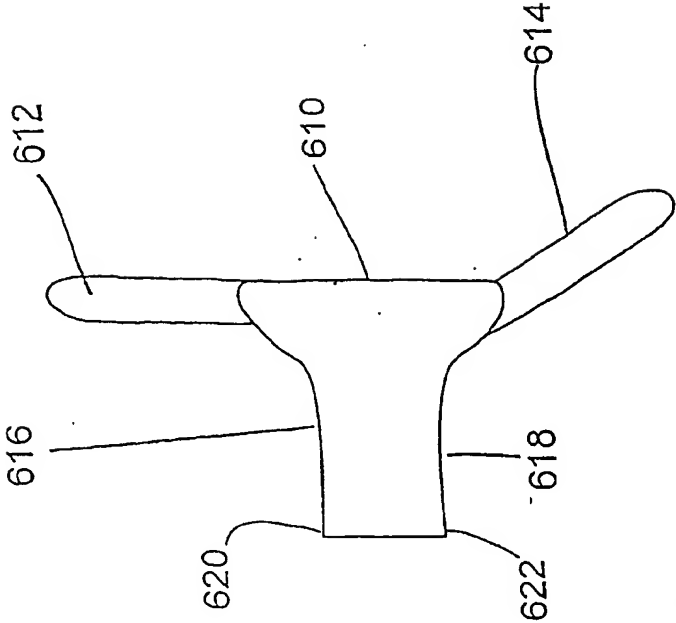


Fig. 71a

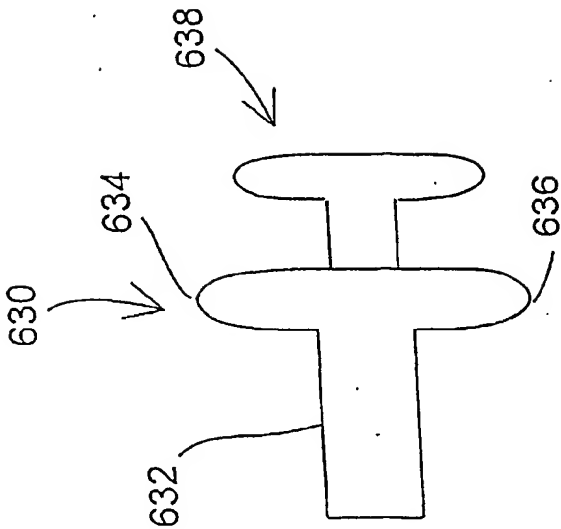


Fig. 72

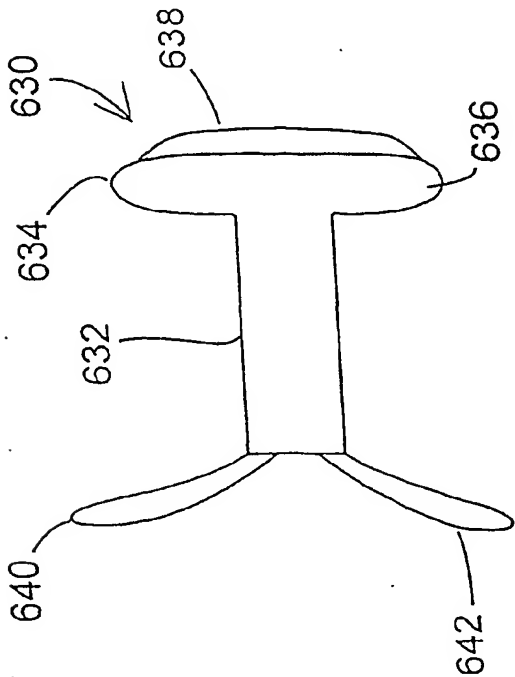


Fig. 73

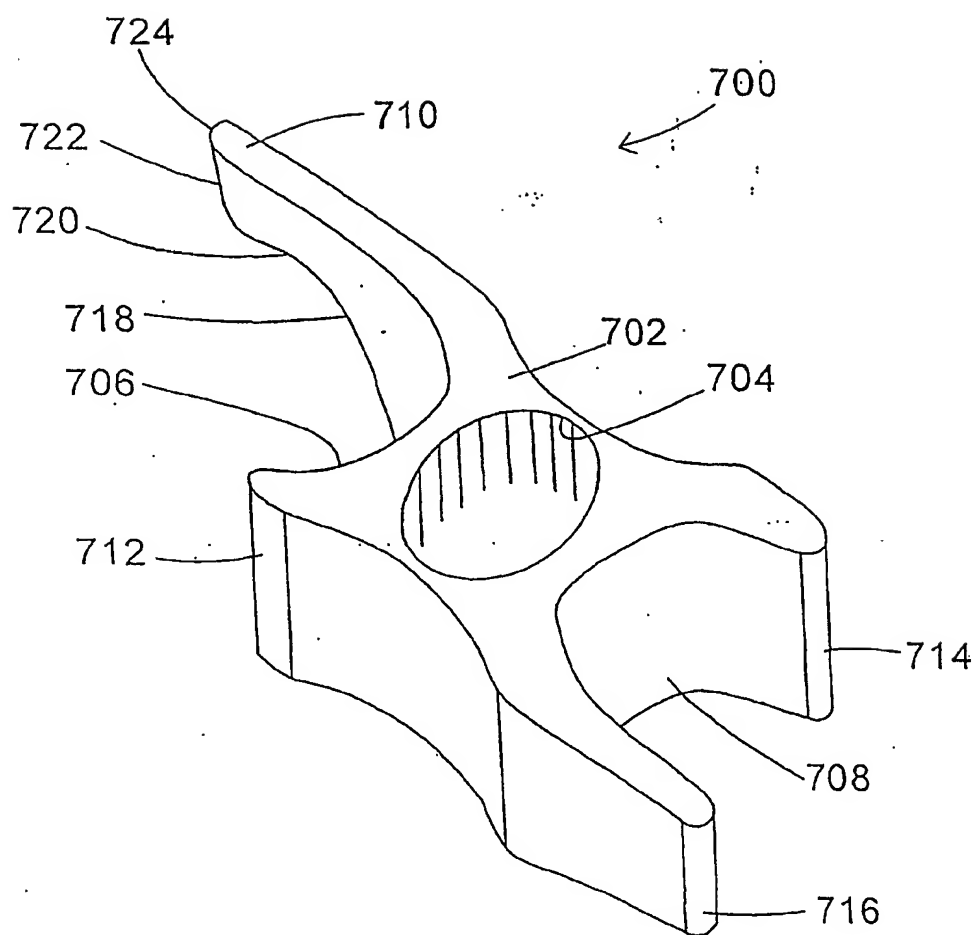


Fig. 74

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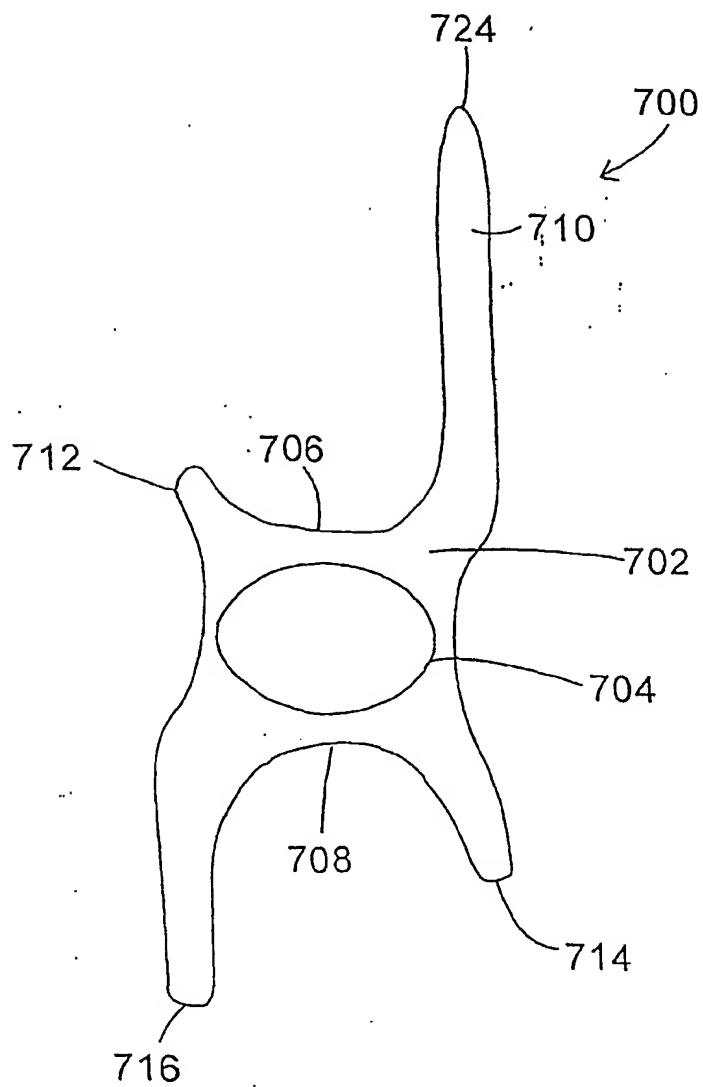
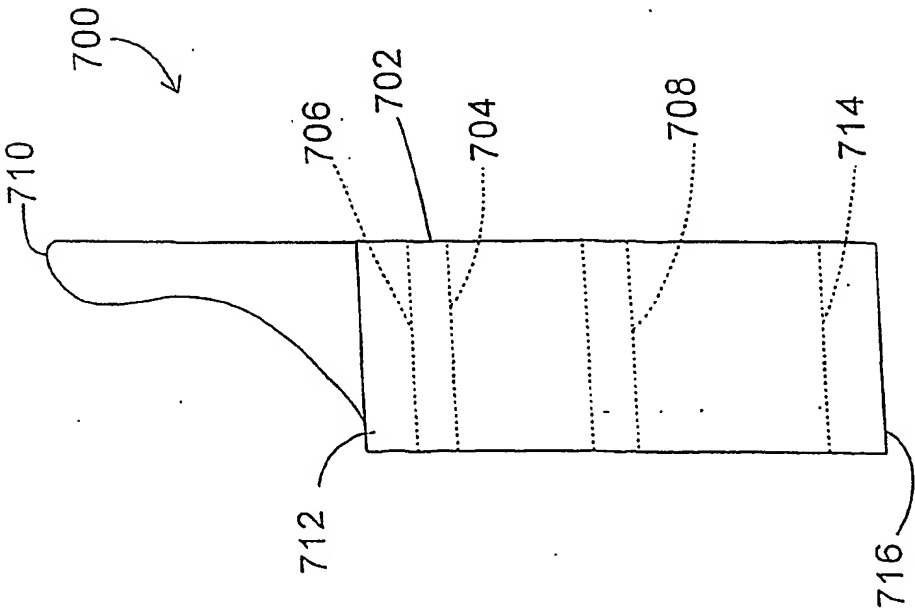
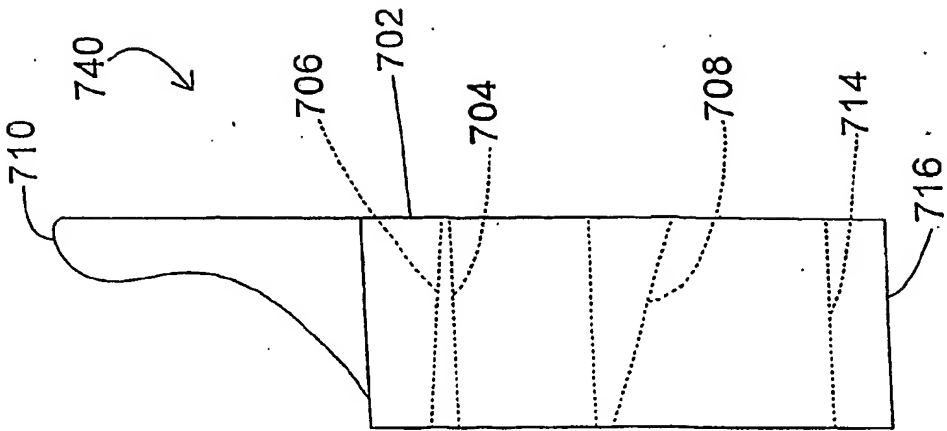


Fig. 75



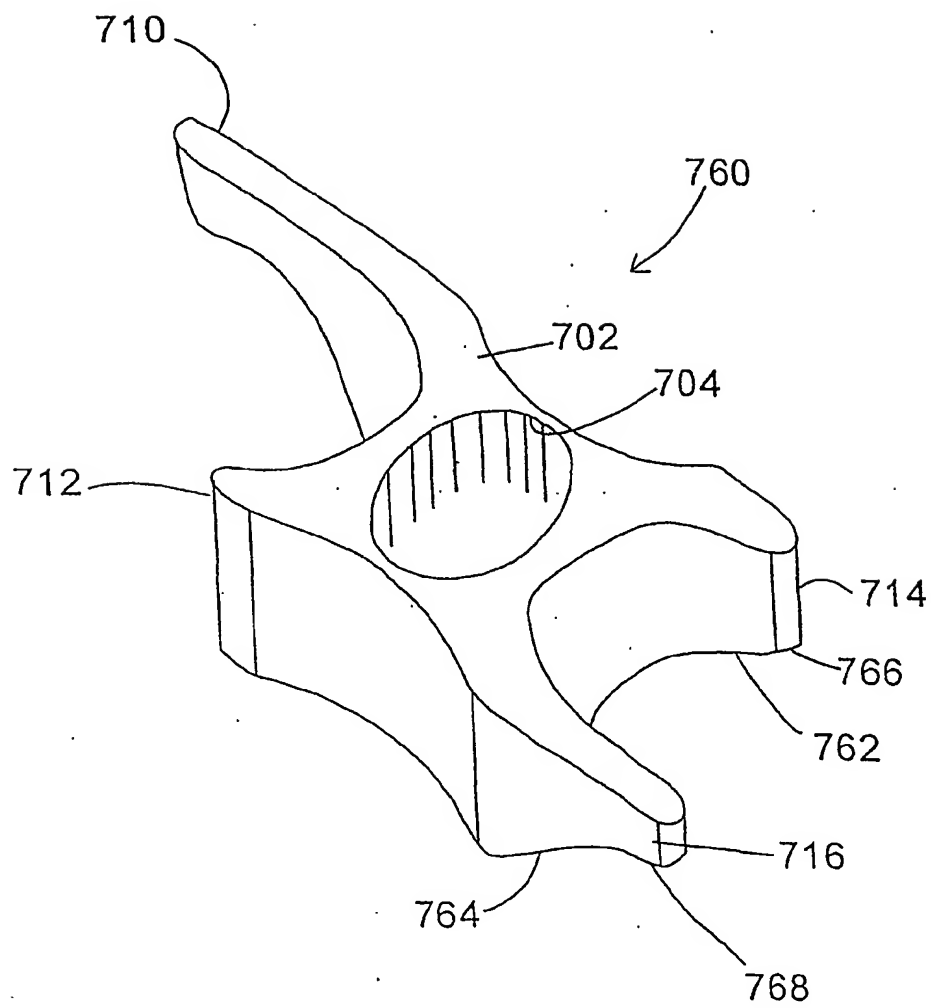


Fig. 78

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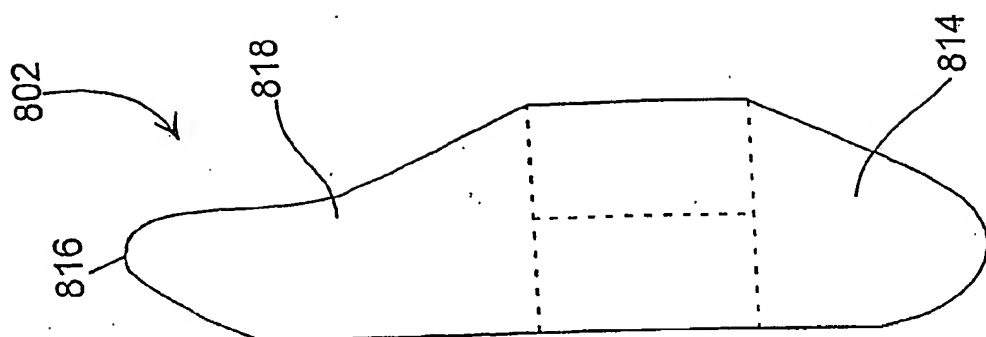


Fig. 81

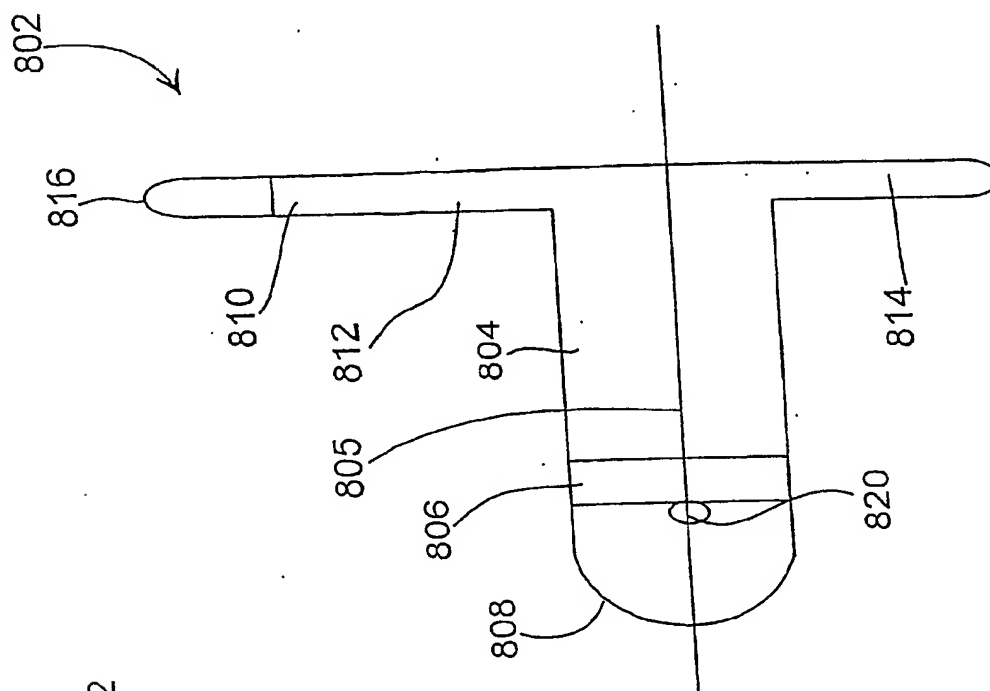


Fig. 80

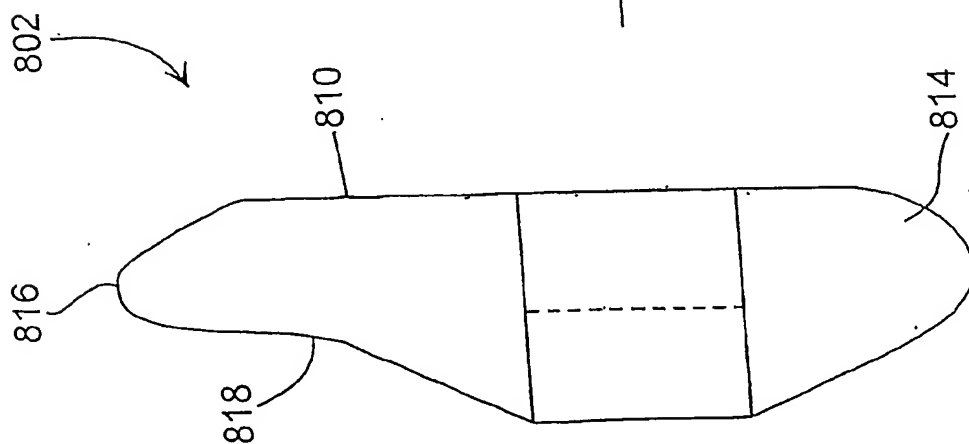
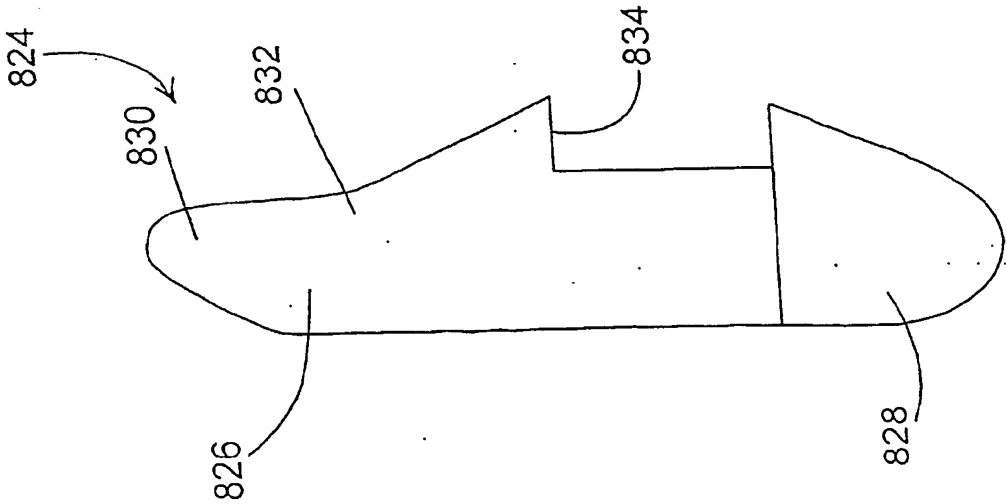
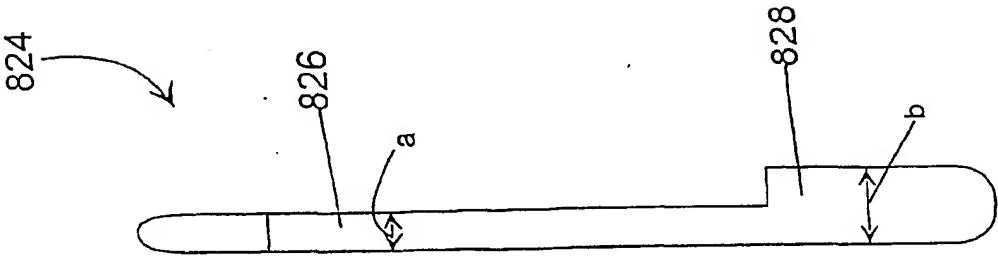
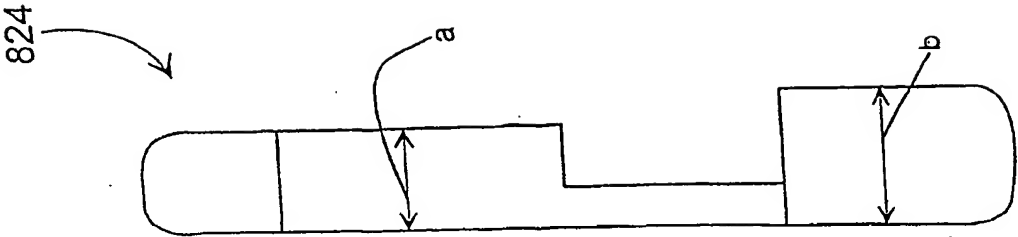


Fig. 79



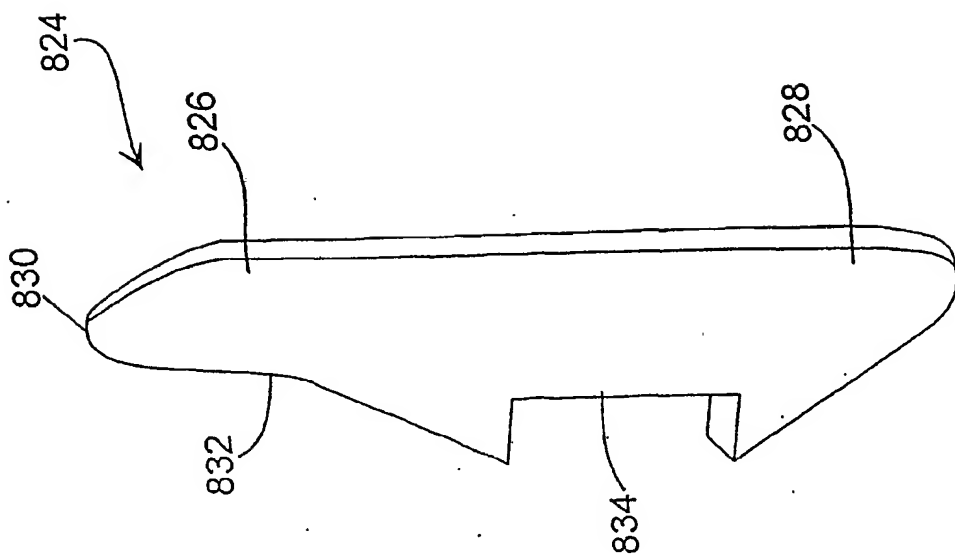


Fig. 85

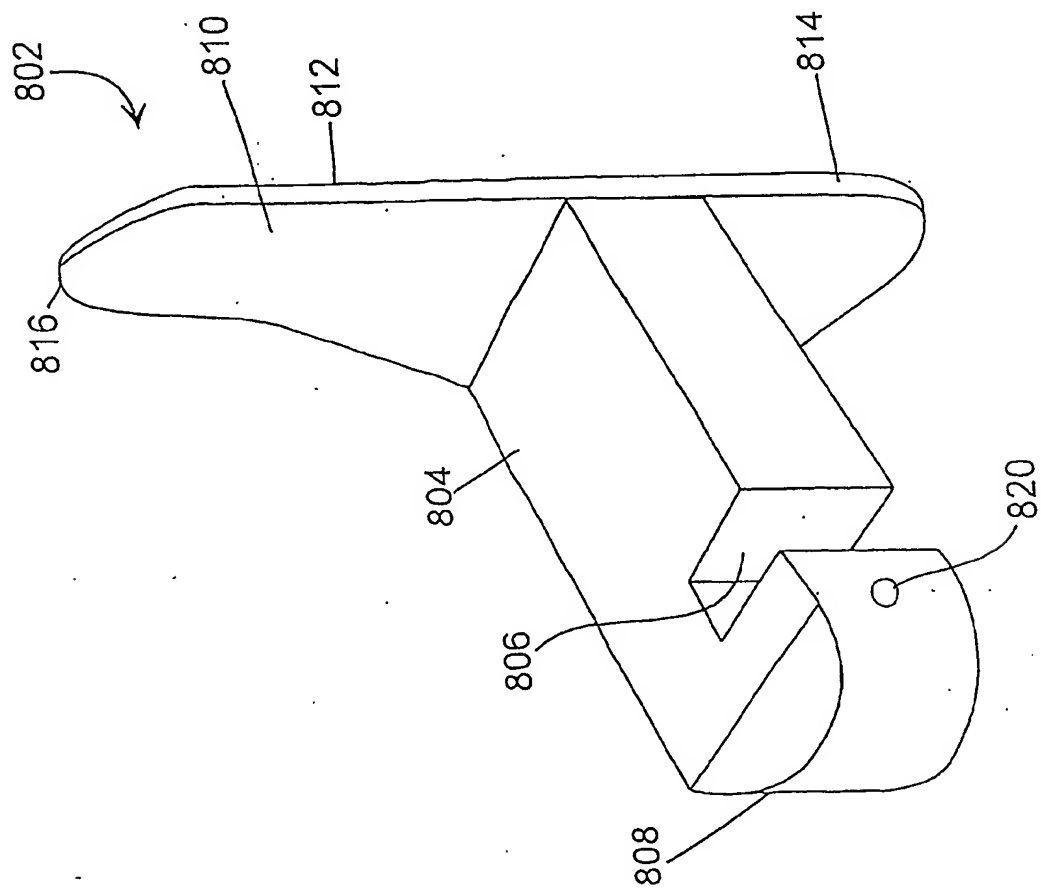


Fig. 84

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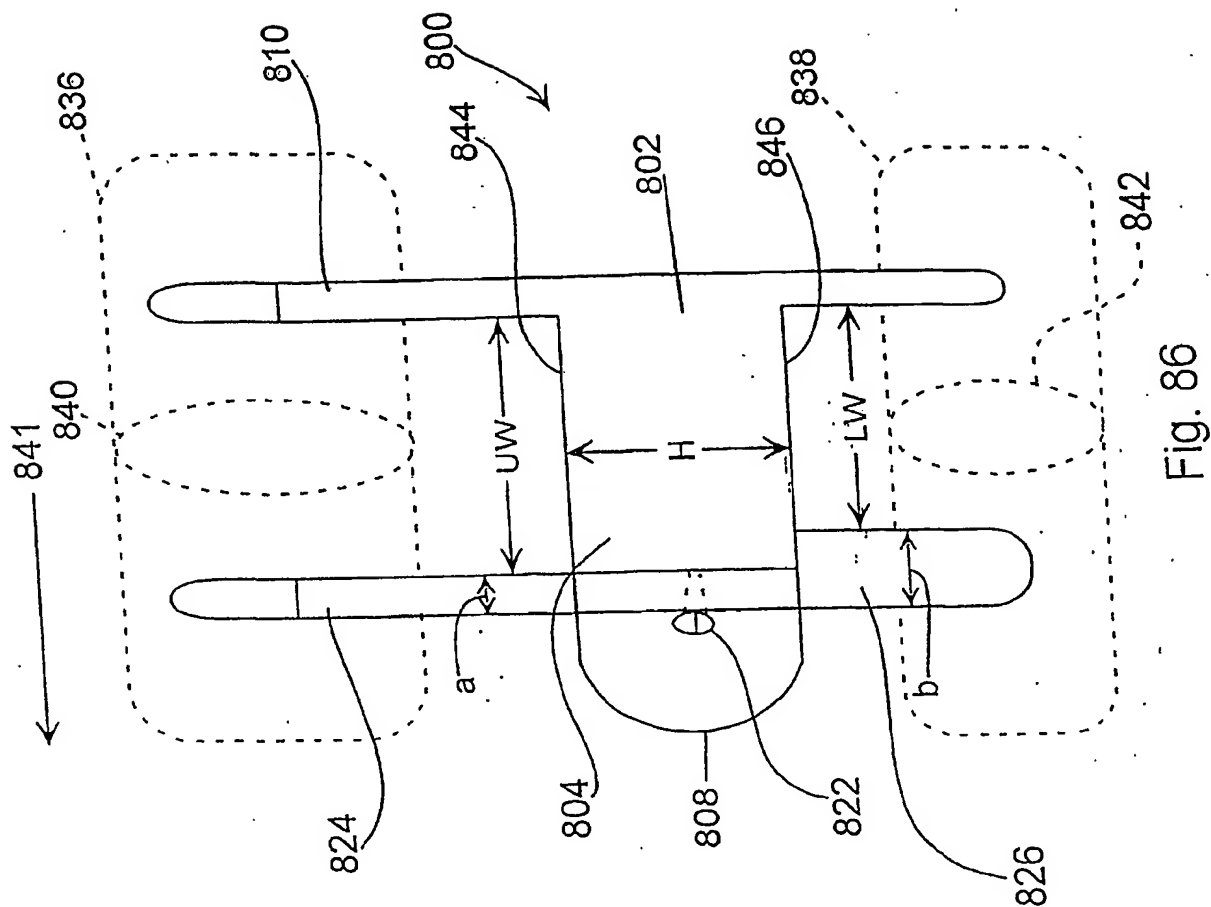
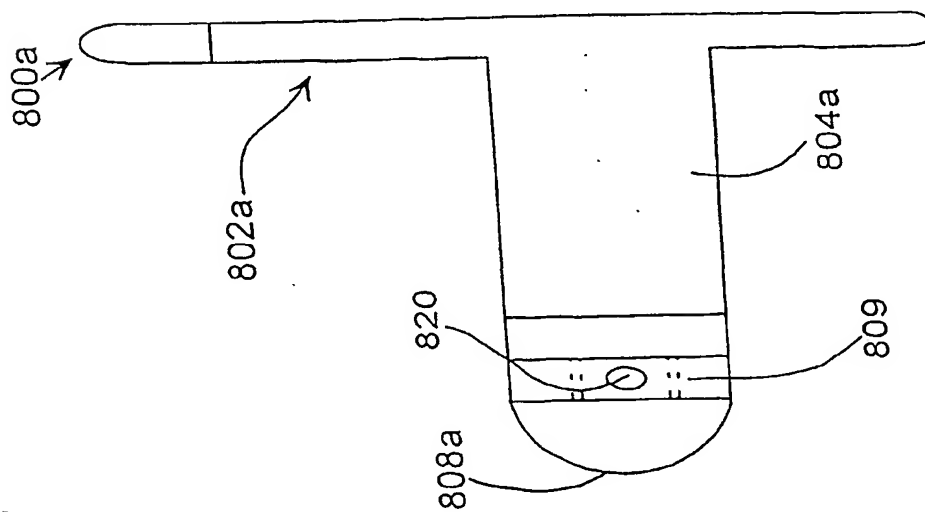


Fig. 80a



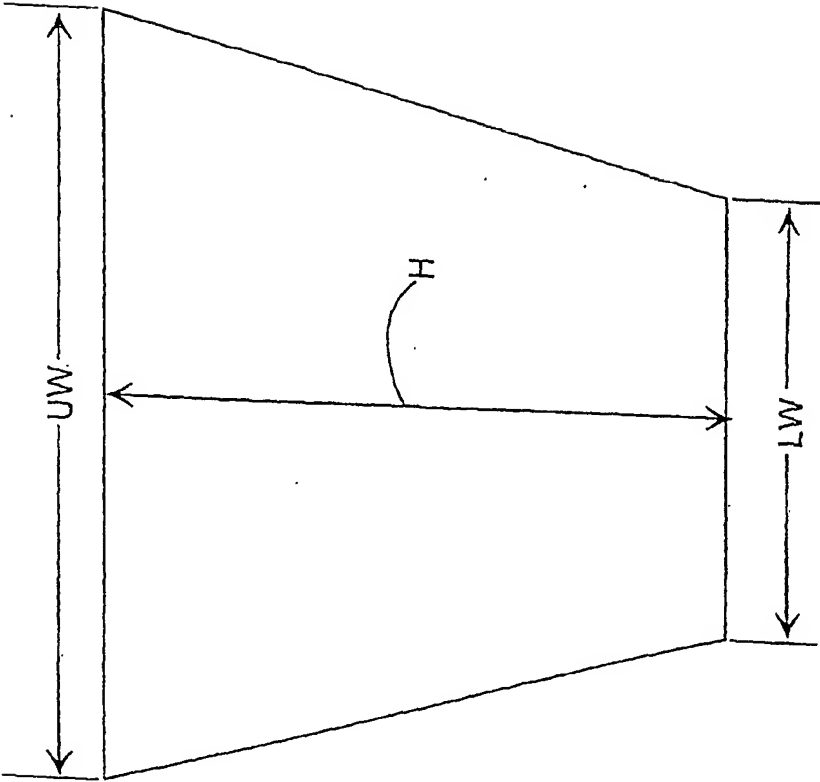
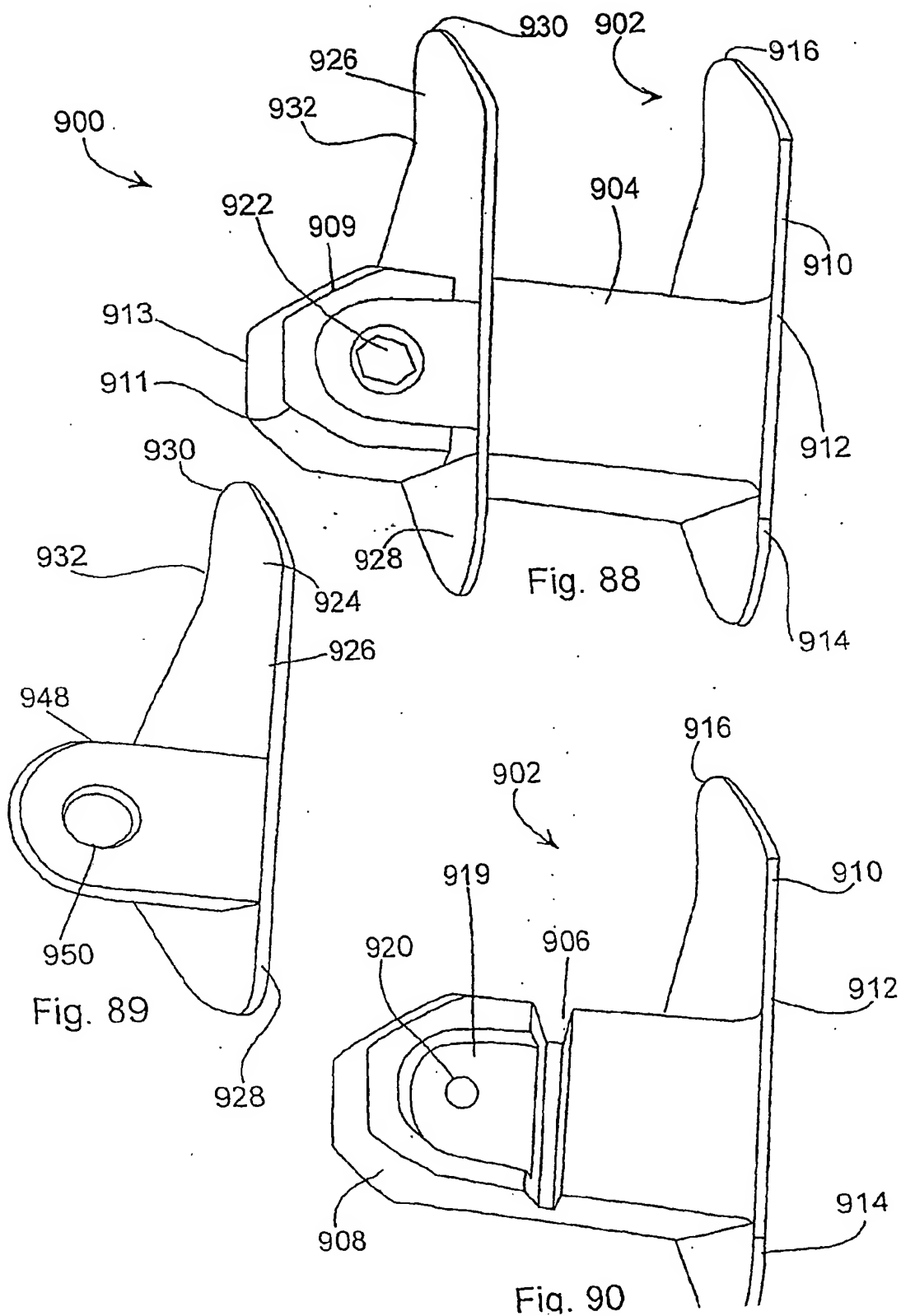


Fig. 87

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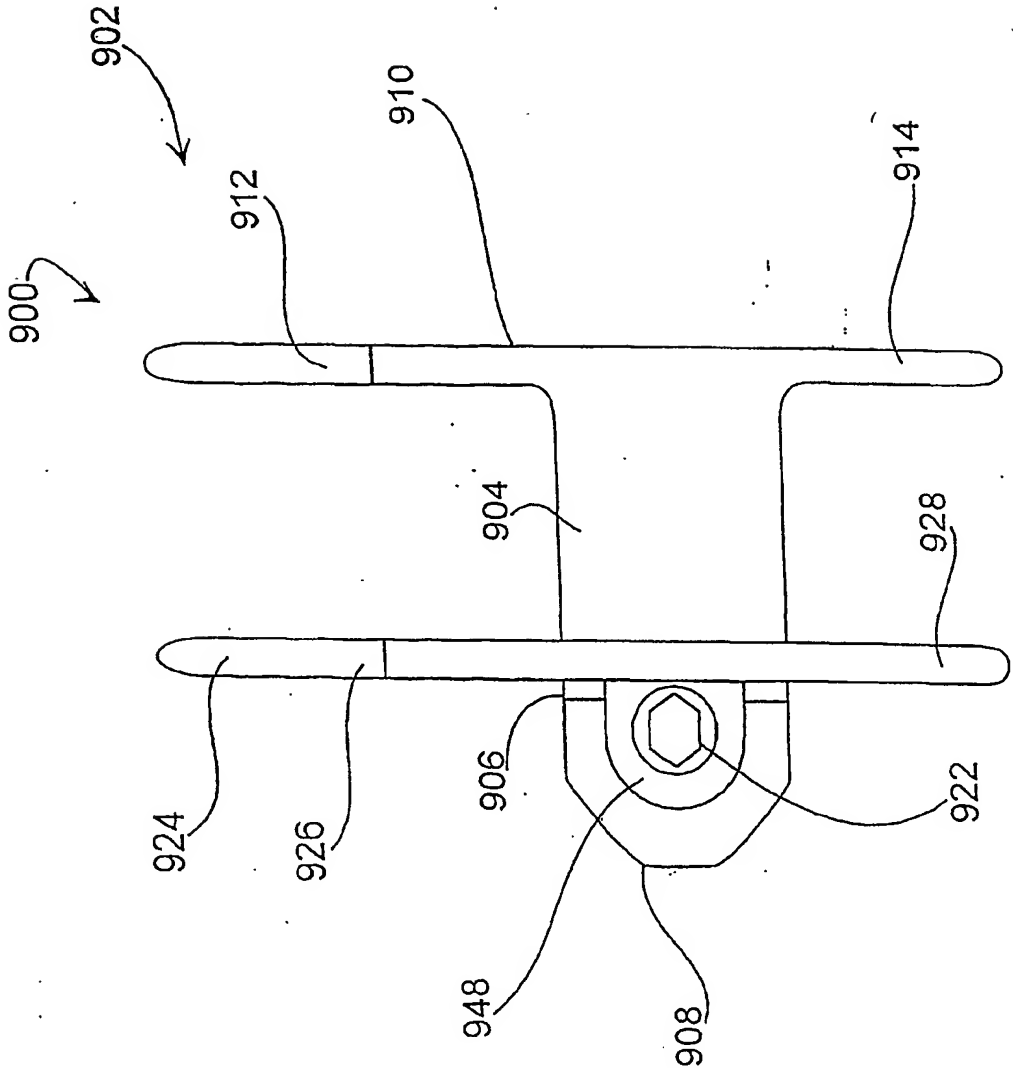


Fig. 91

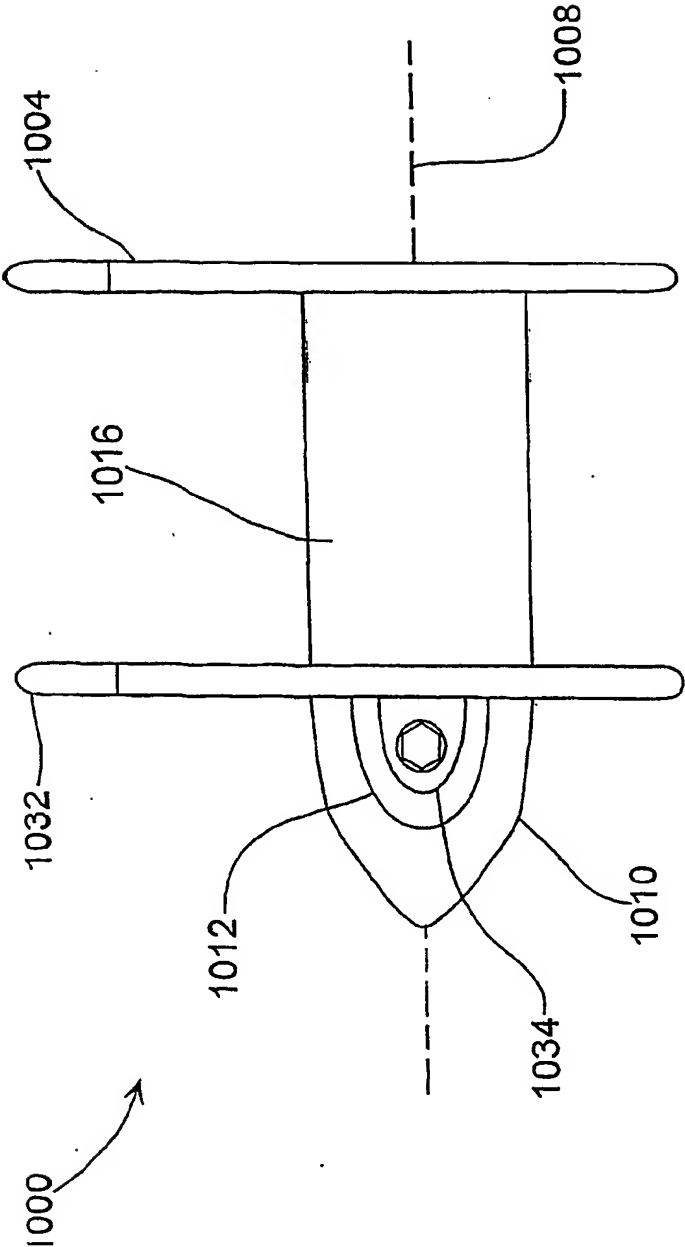


Fig. 92

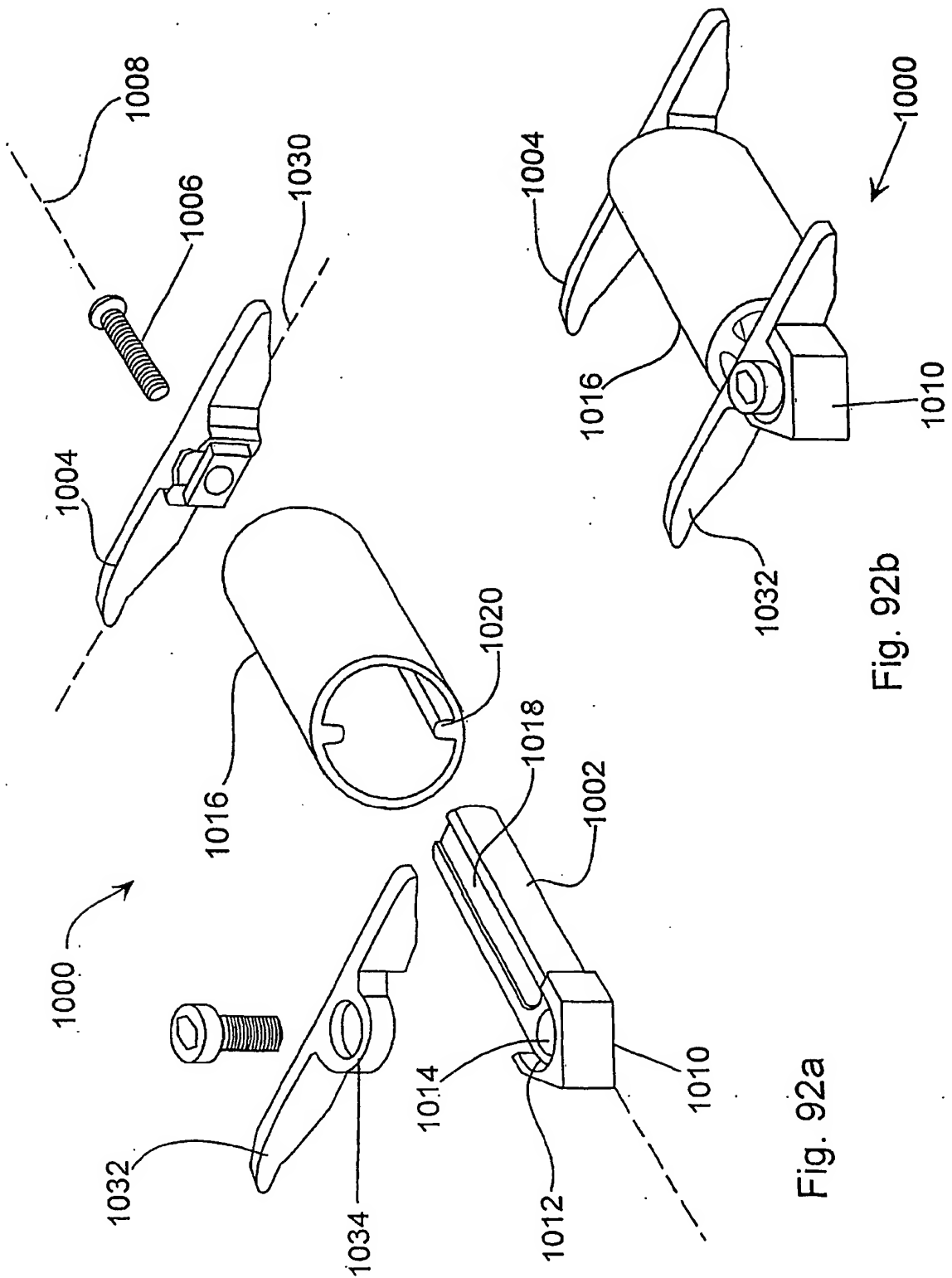
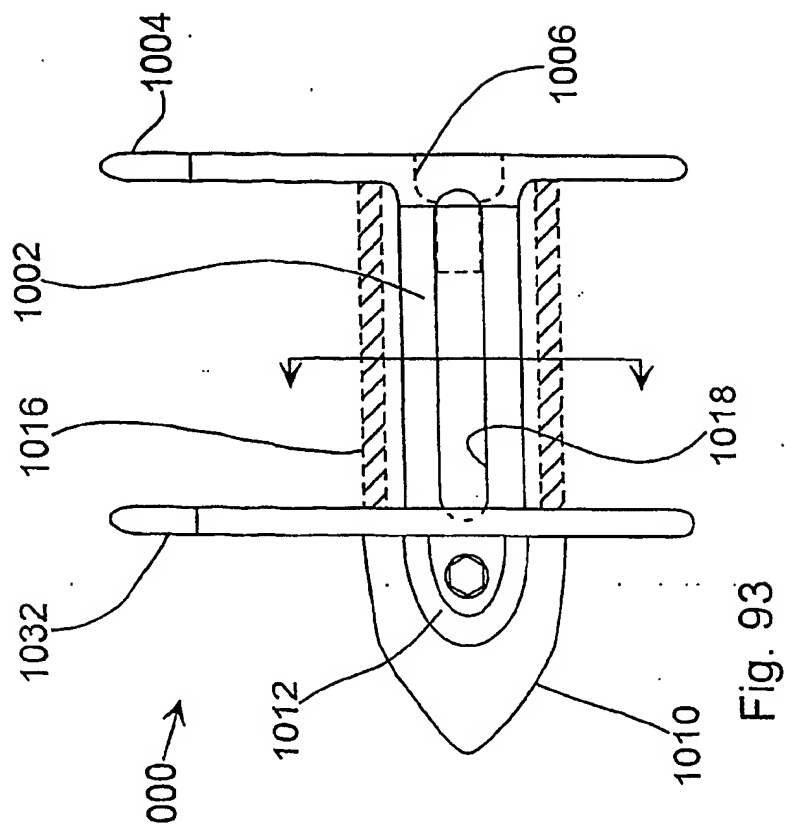
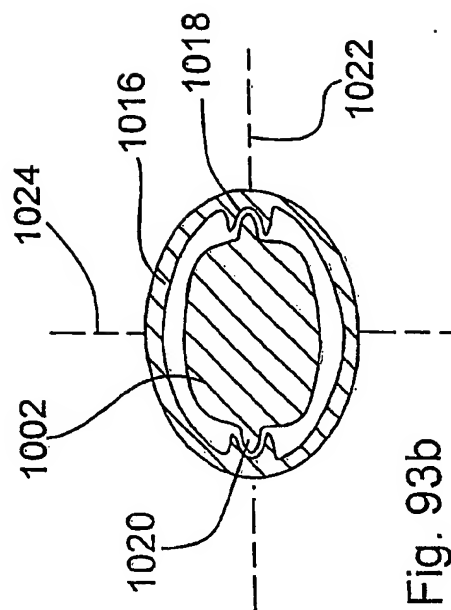
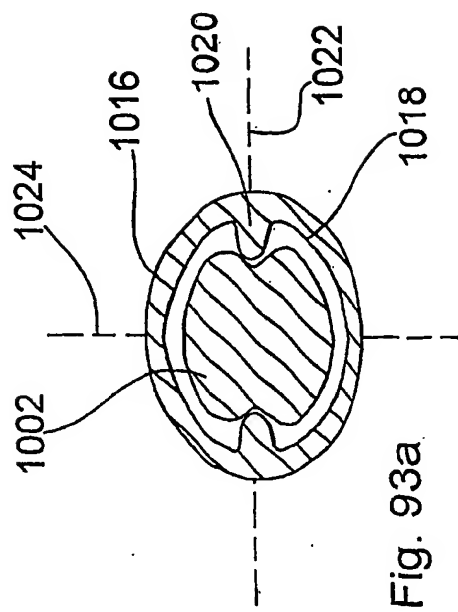


Fig. 92a

Fig. 92b



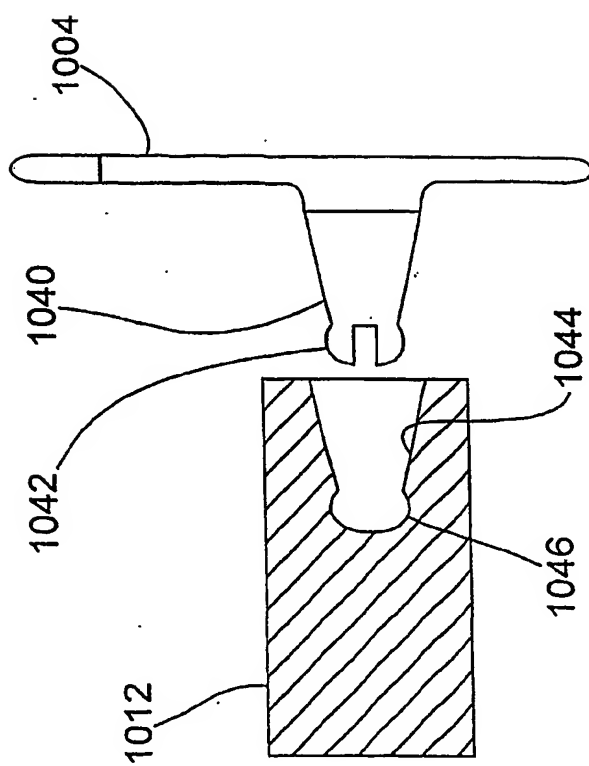


Fig. 93c

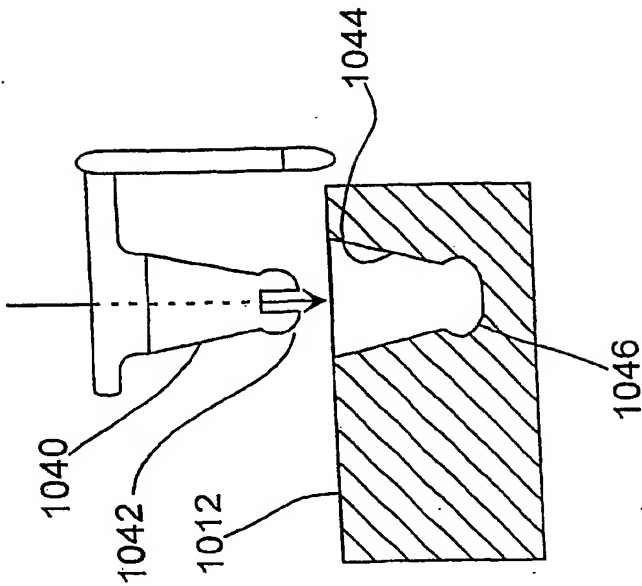


Fig. 93d

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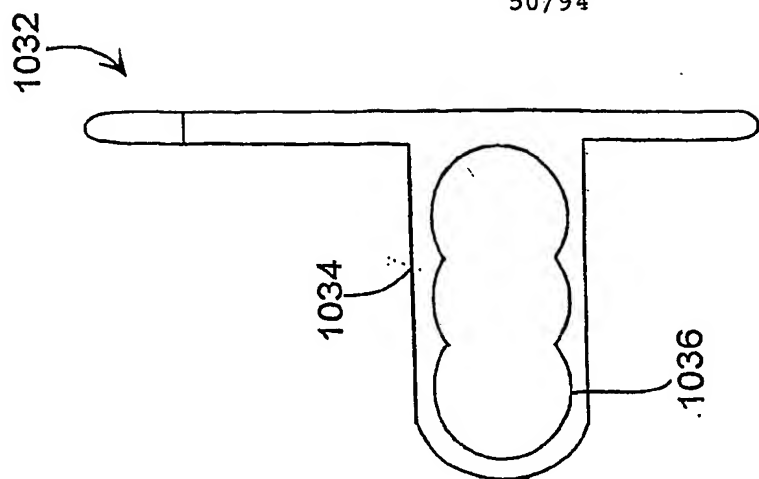


Fig. 94b

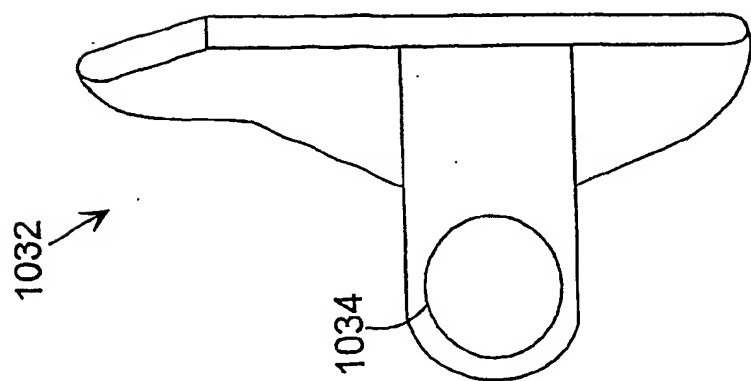


Fig. 94a

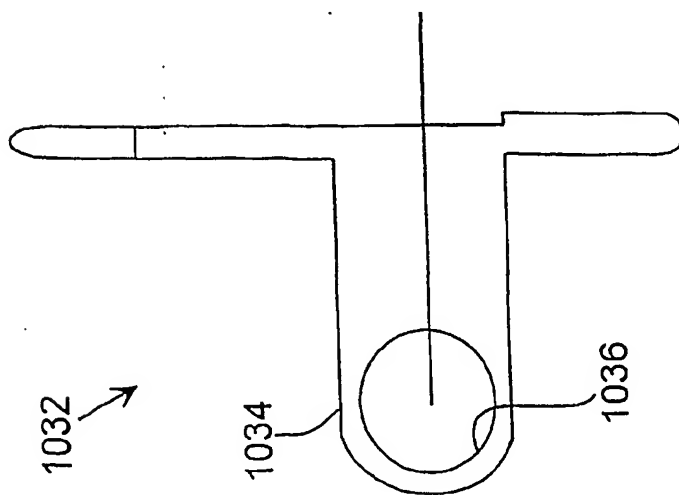


Fig. 94

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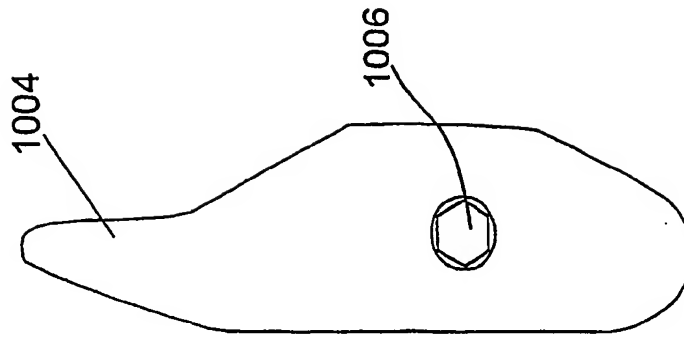


Fig. 96

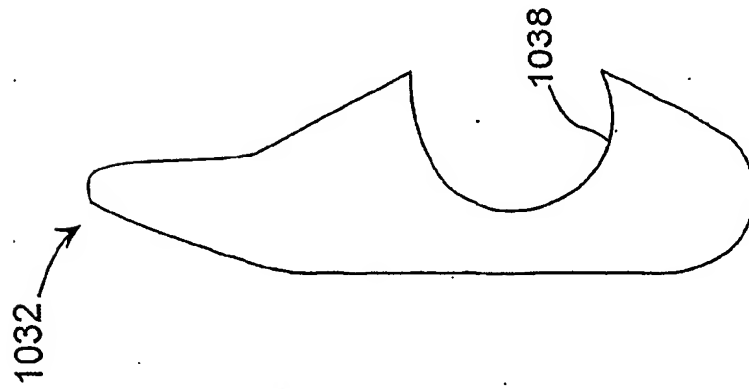


Fig. 95a

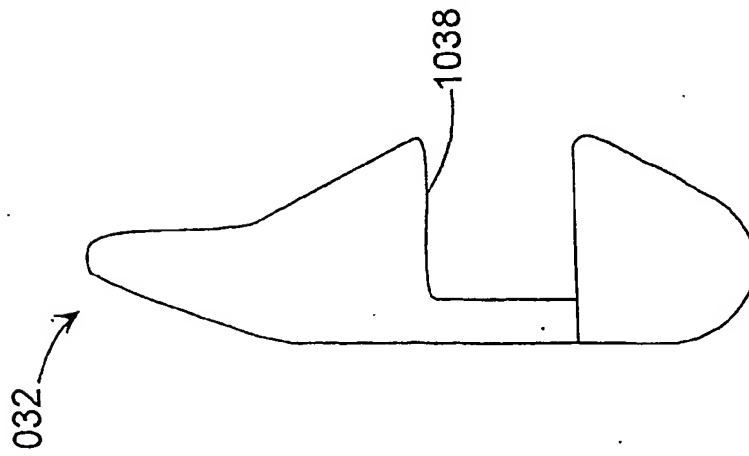


Fig. 95

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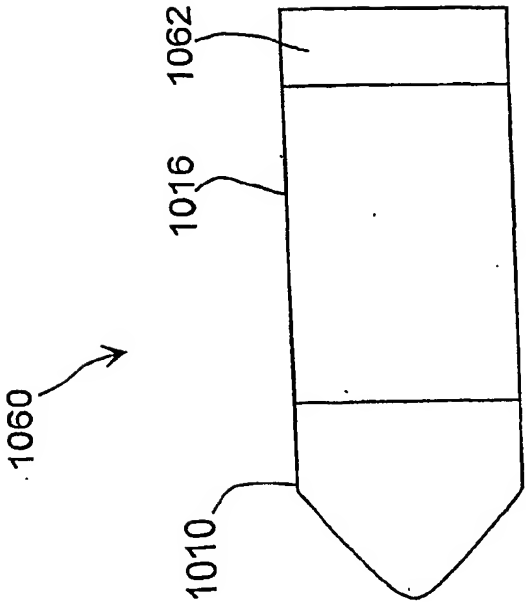


Fig. 98

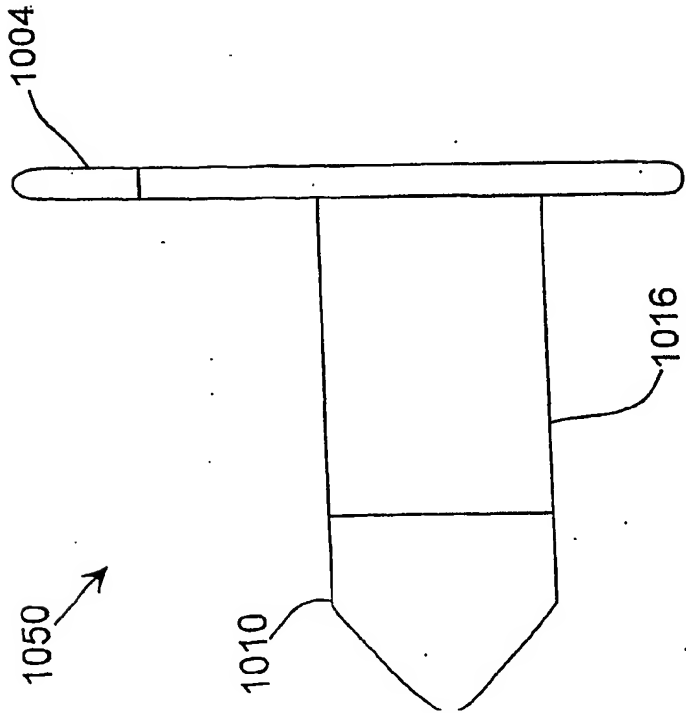


Fig. 97

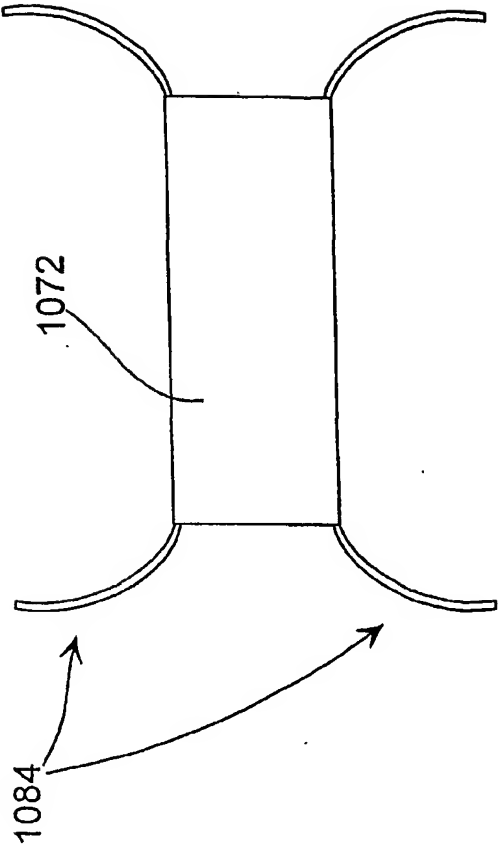


Fig. 99

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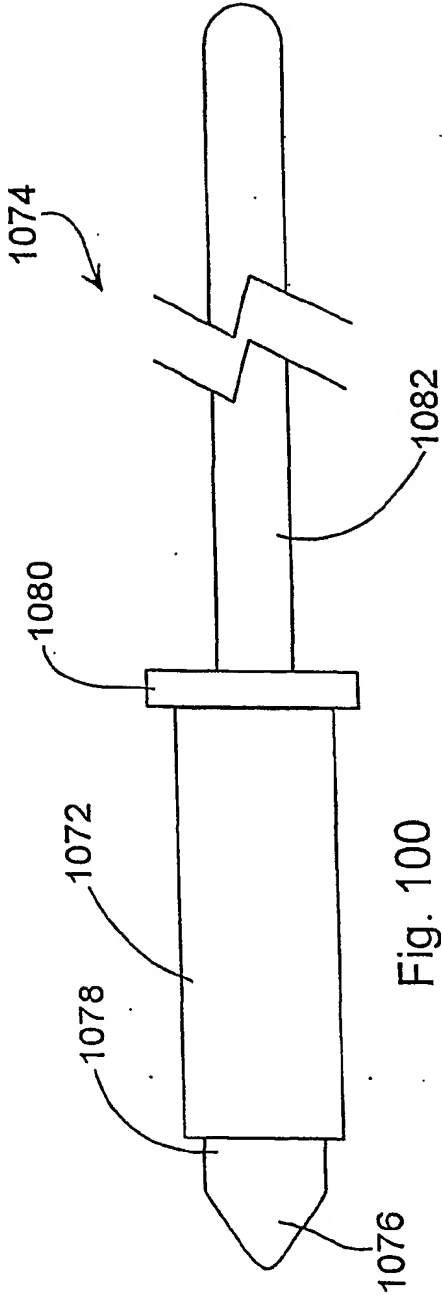


Fig. 100

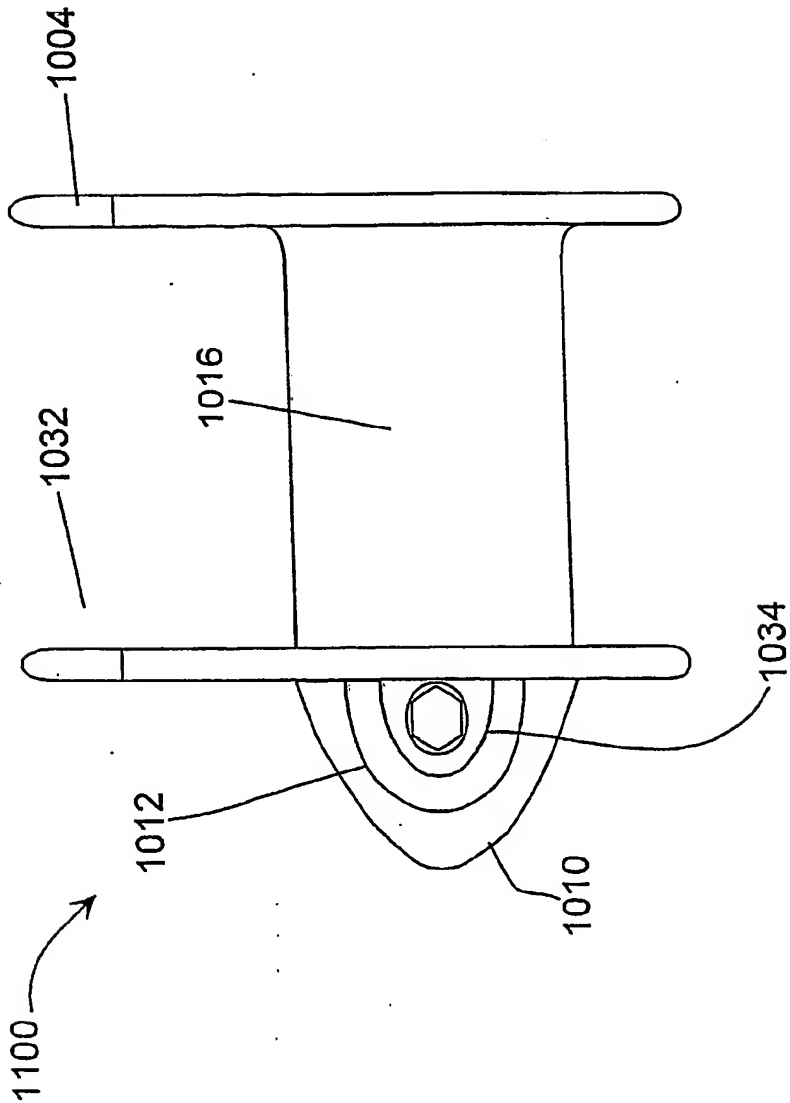


Fig. 101

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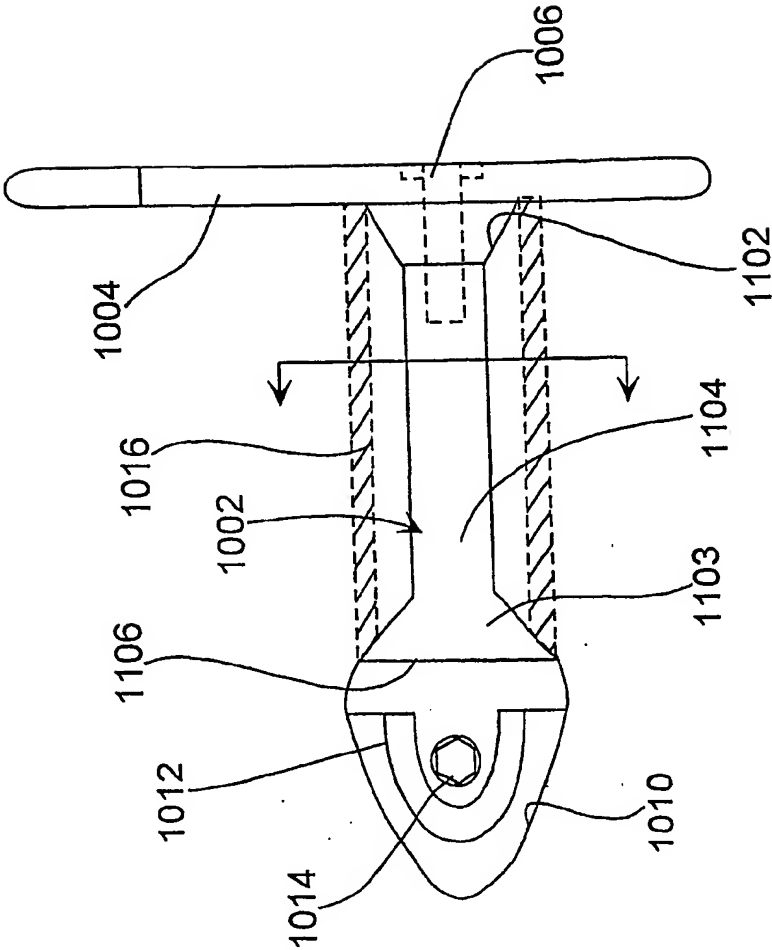


Fig. 102

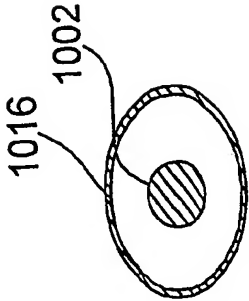


Fig. 102a

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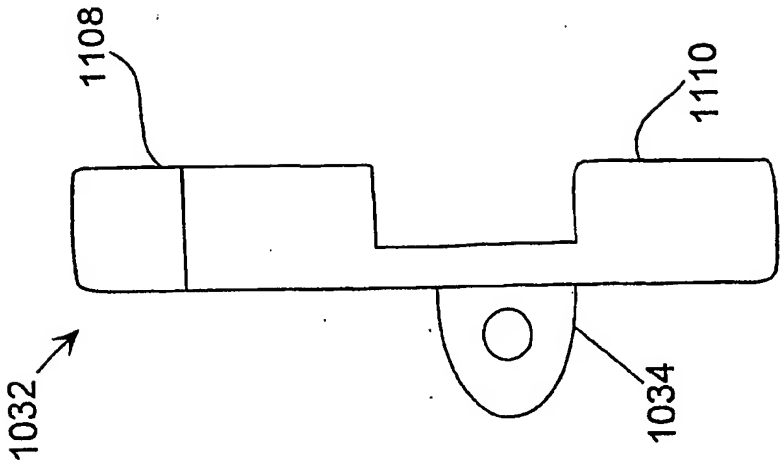


Fig. 103

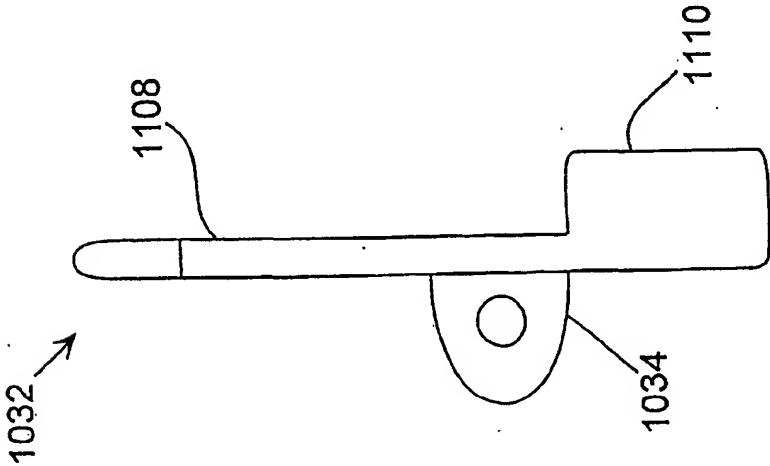


Fig. 104

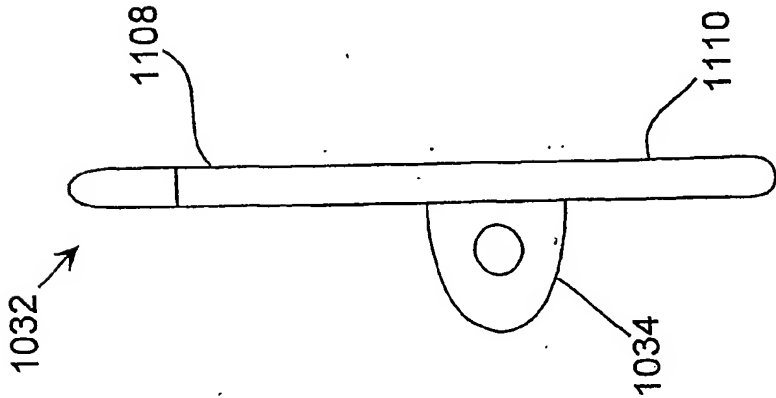


Fig. 105

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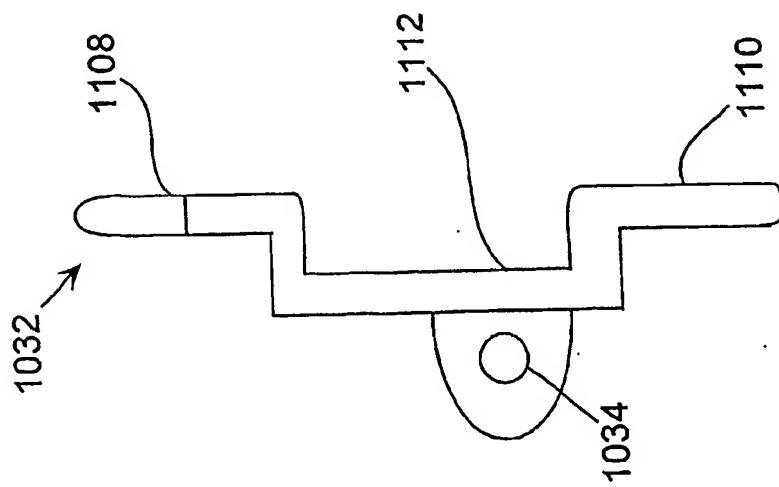


Fig. 107

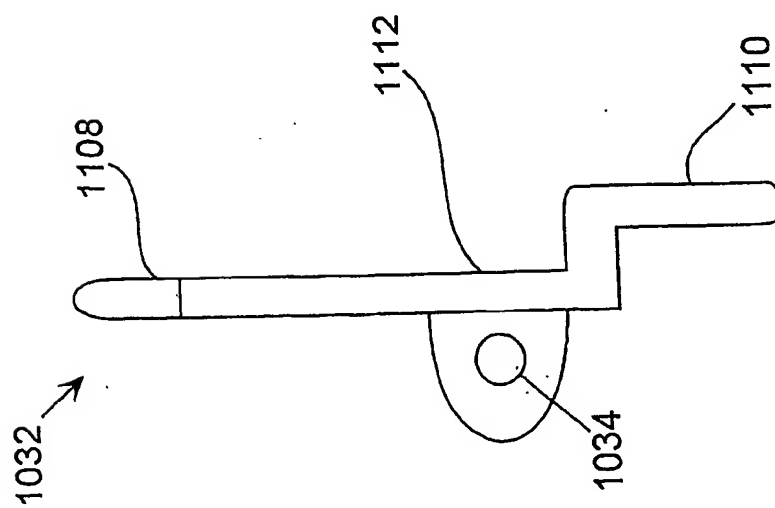


Fig. 106

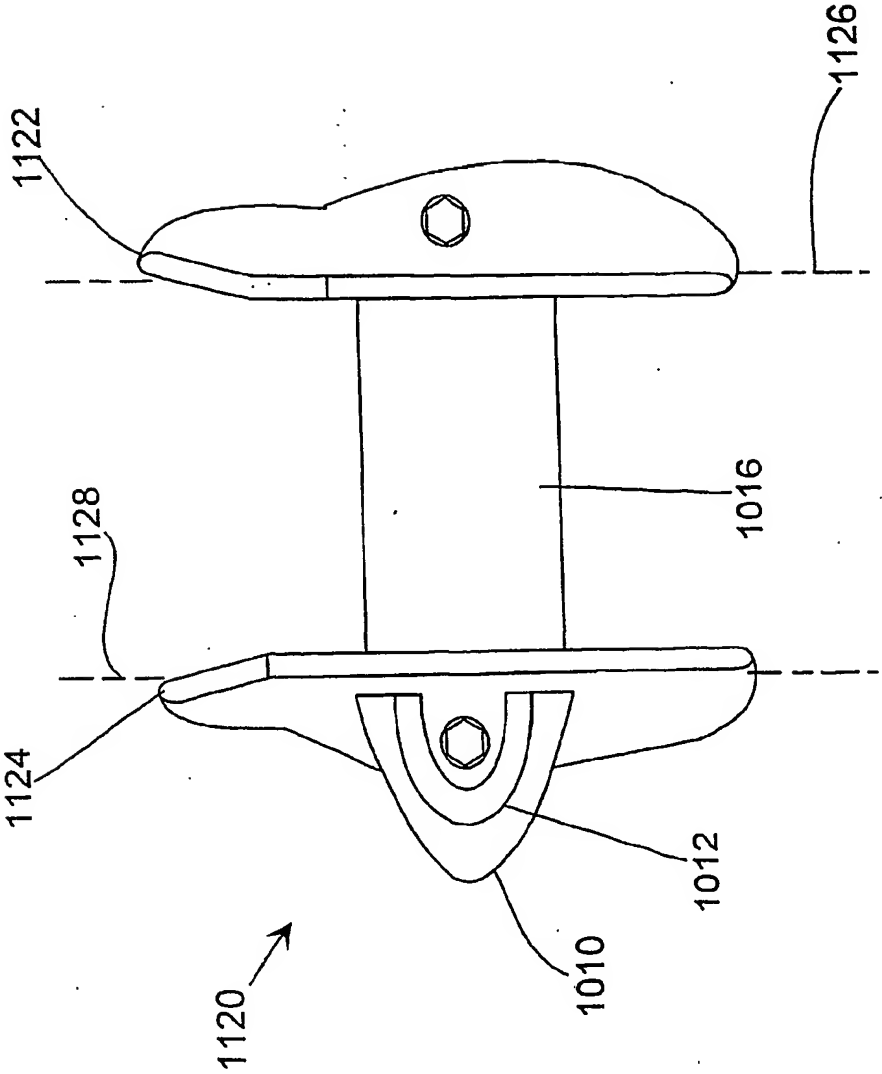
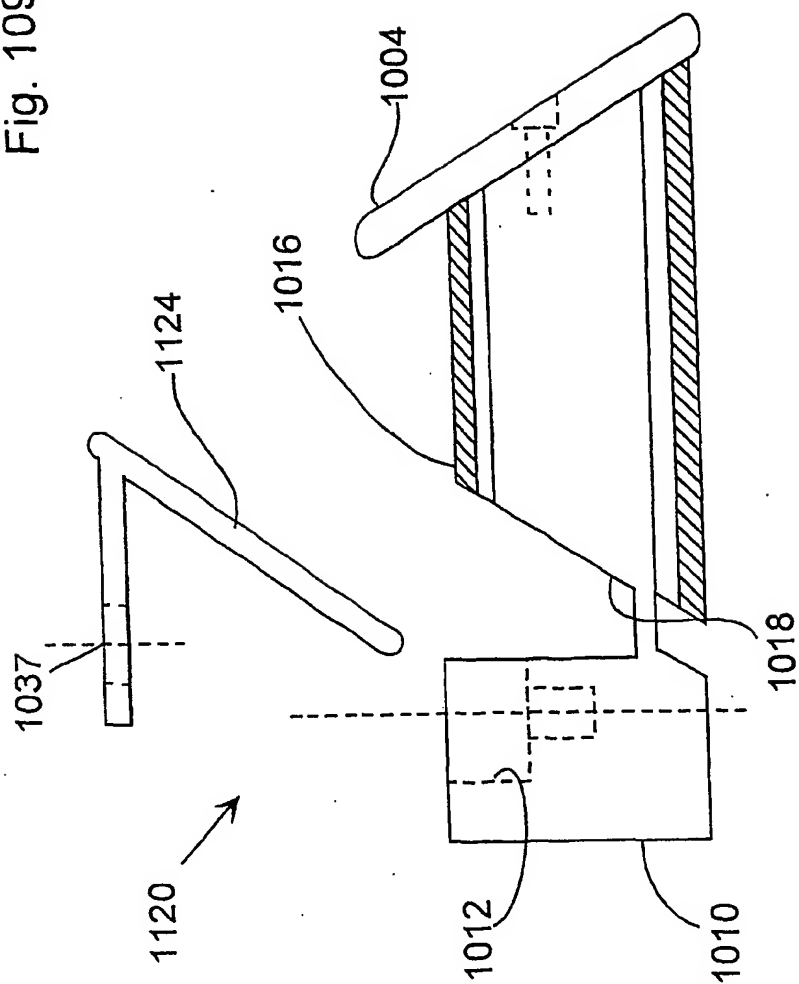


Fig. 108

Fig. 109



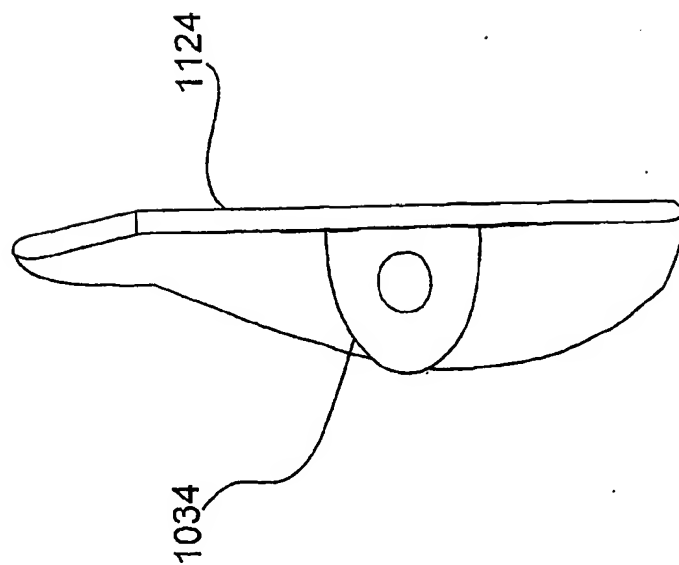
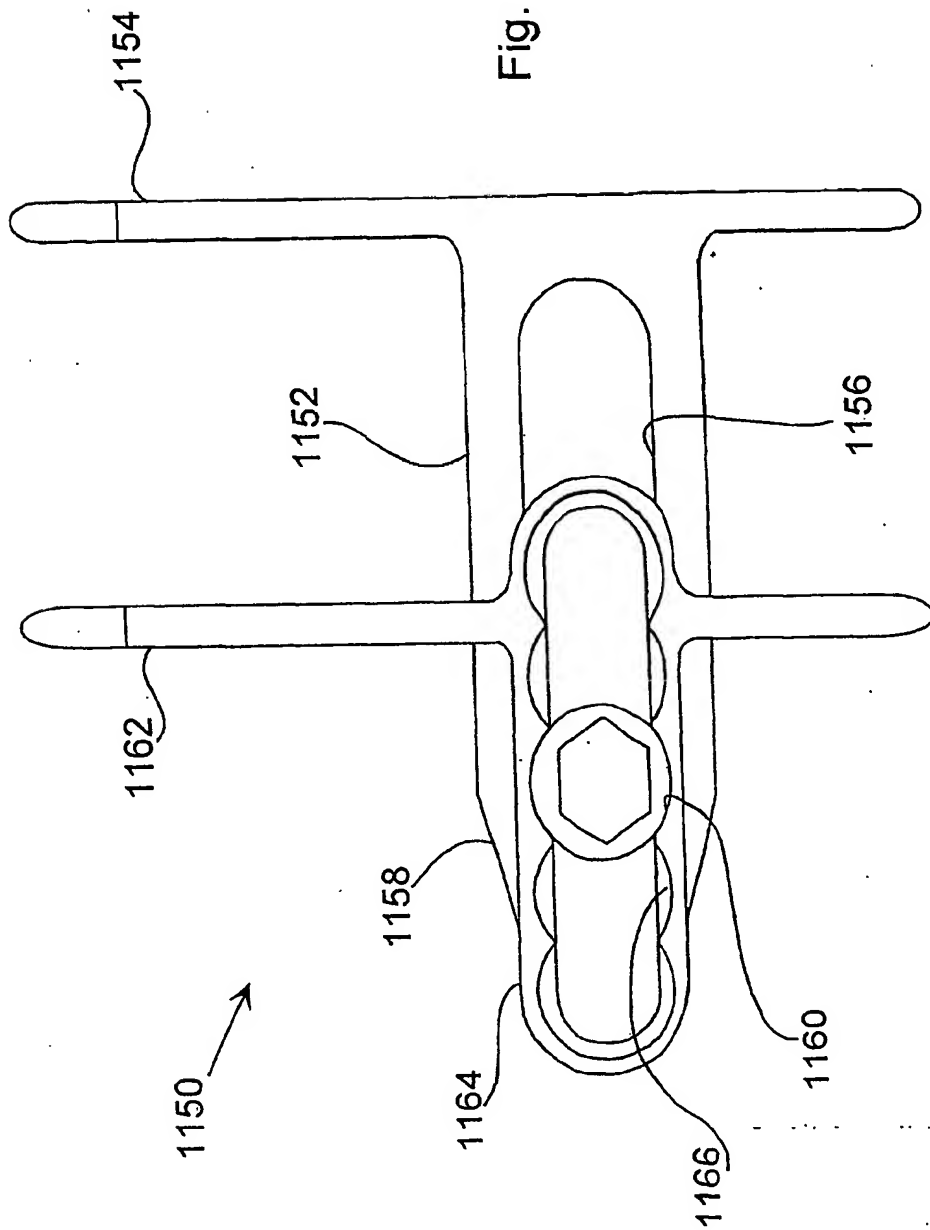


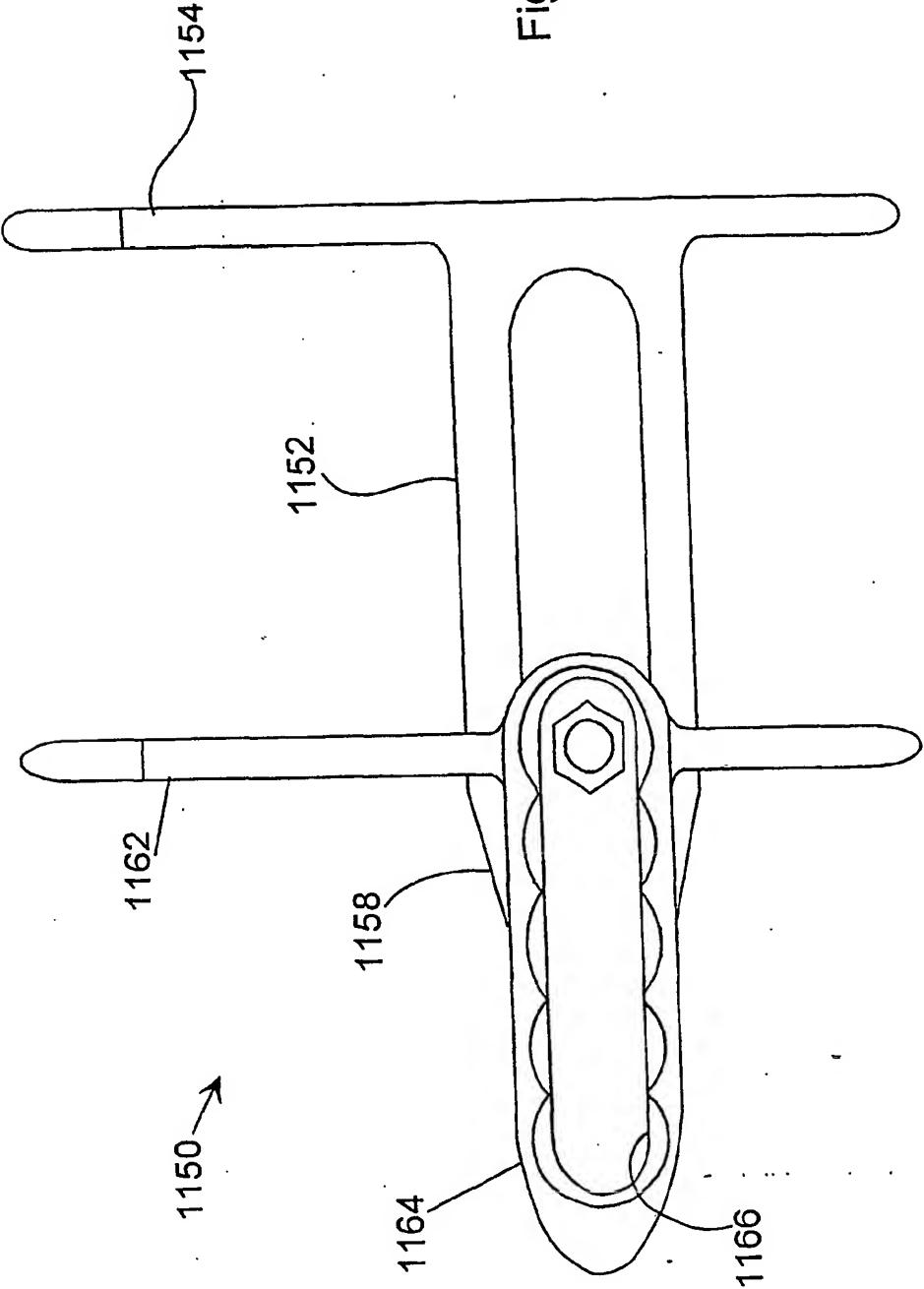
Fig. 110

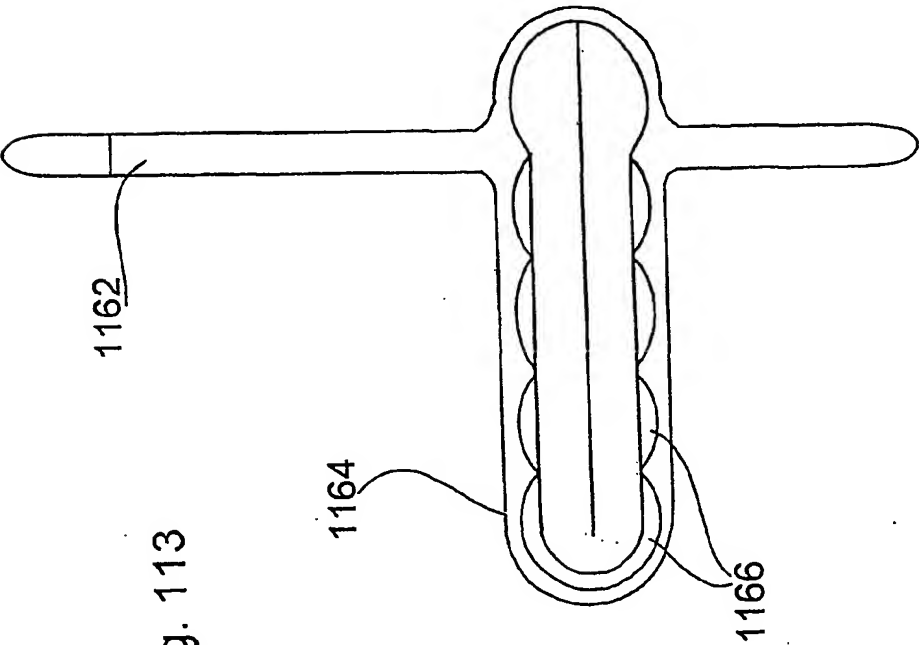
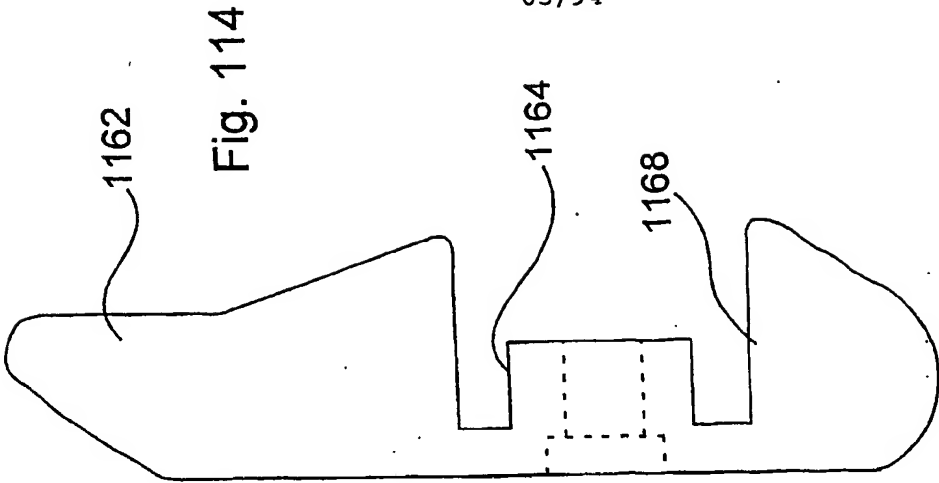
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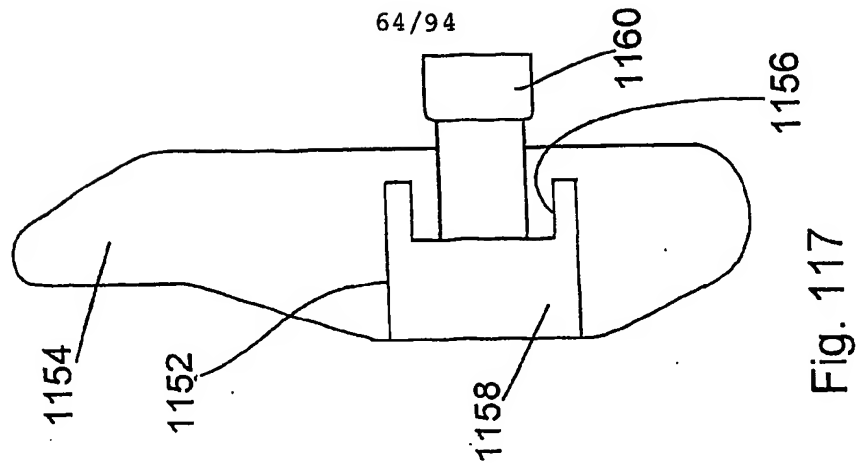
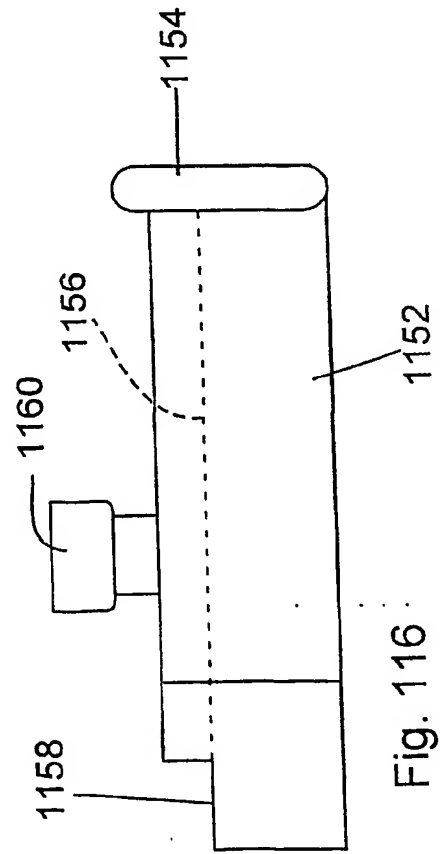
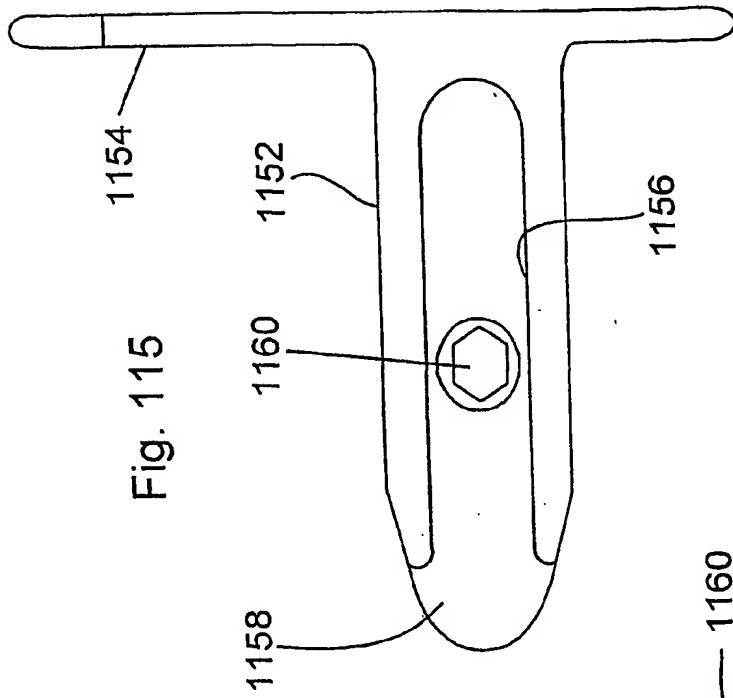
Fig. 111



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Fig. 112

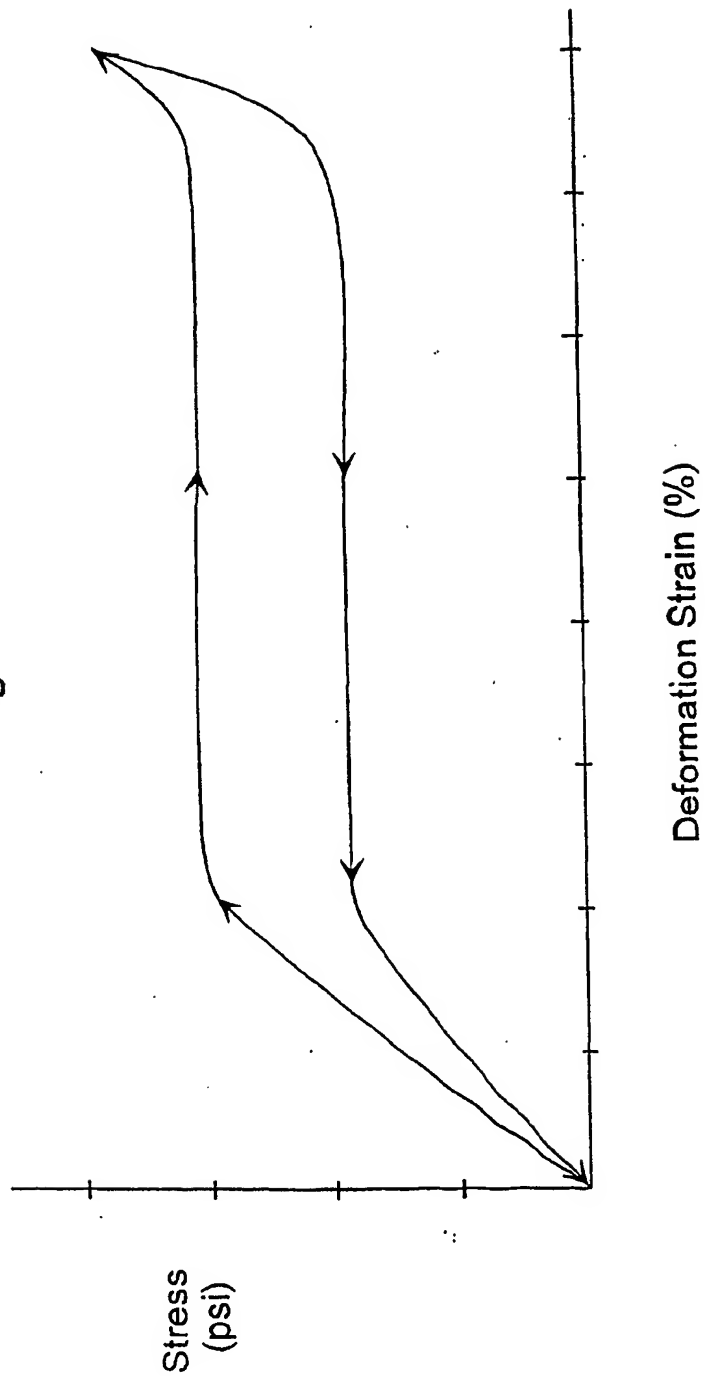






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Fig. 118



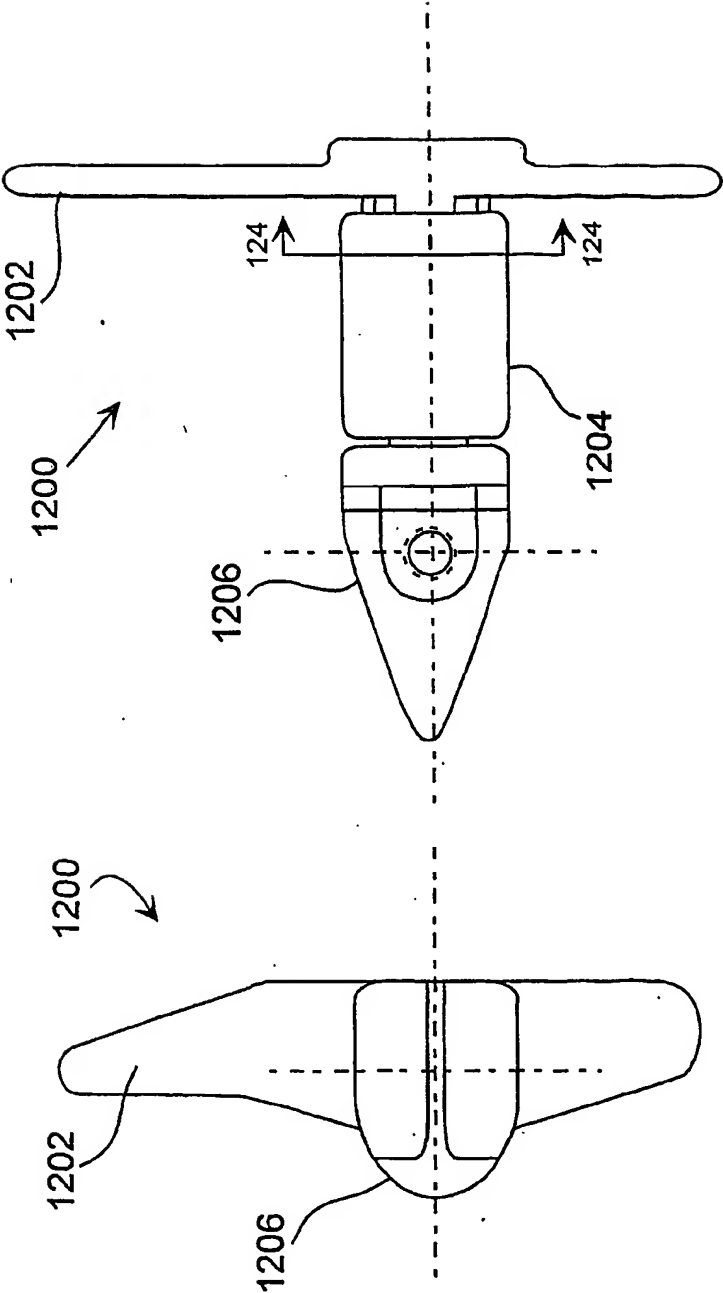


Fig. 119b

Fig. 119a

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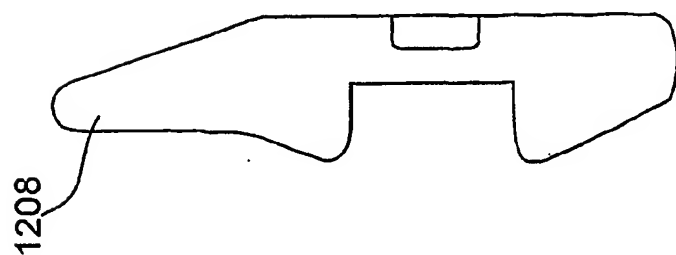


Fig. 120a

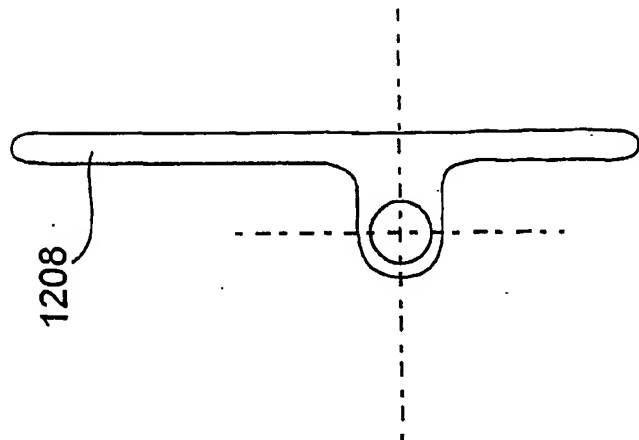


Fig. 120b

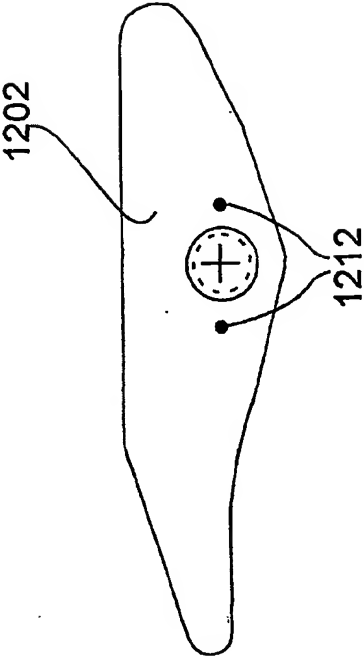


Fig. 121a

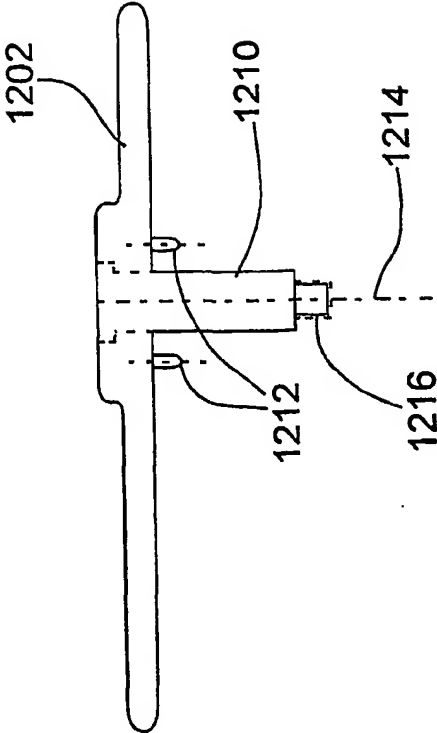


Fig. 121b

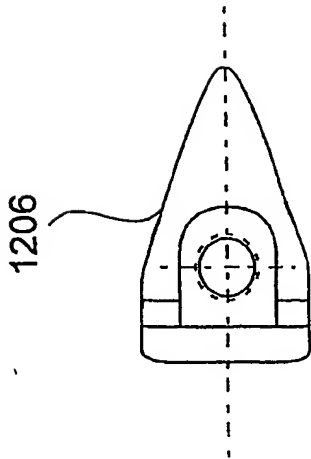


Fig. 122a

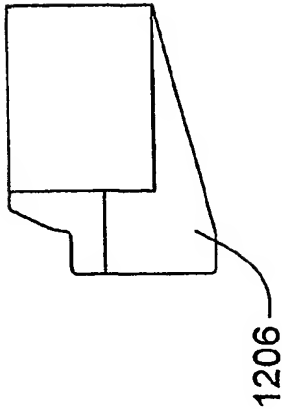


Fig. 122b

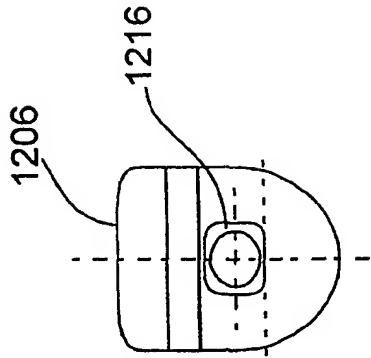


Fig. 122c

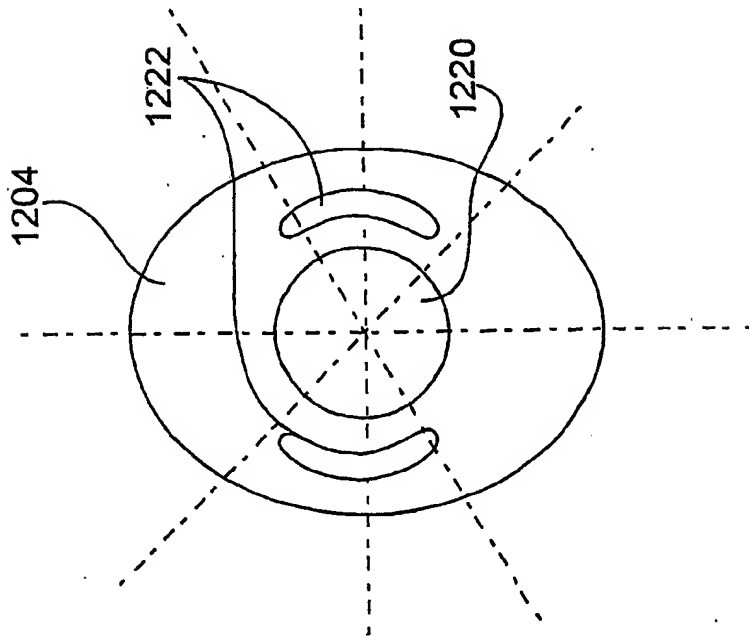


Fig. 123a

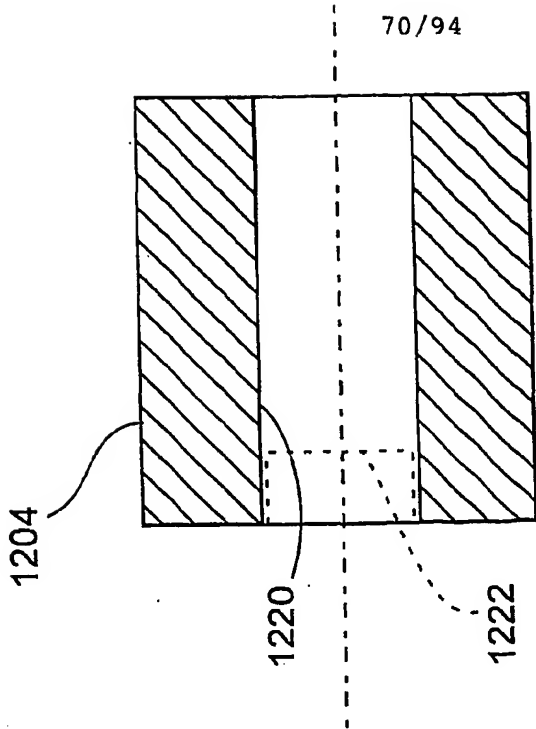


Fig. 123b

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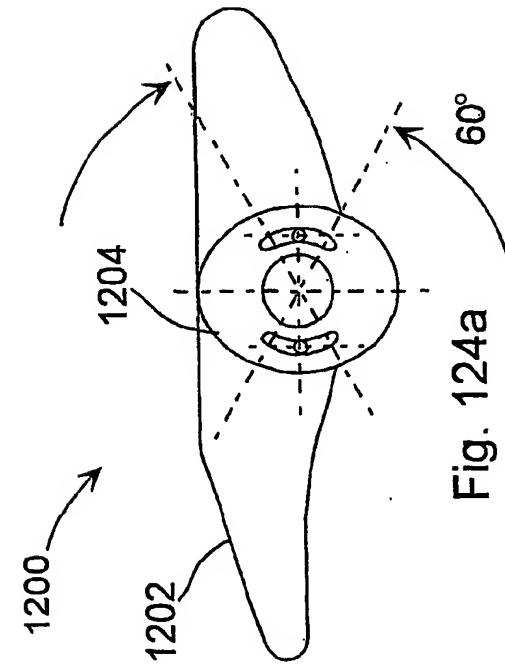


Fig. 124a

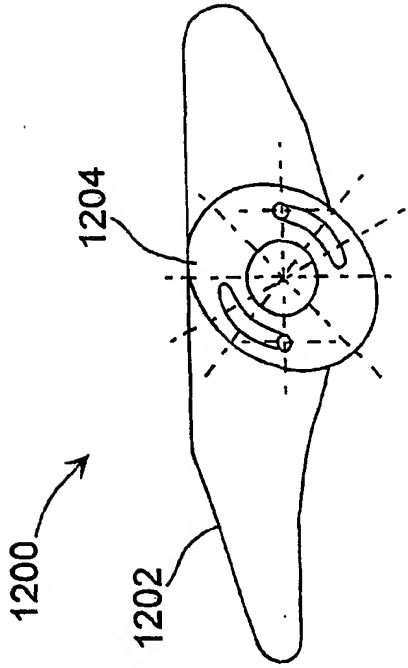


Fig. 124c

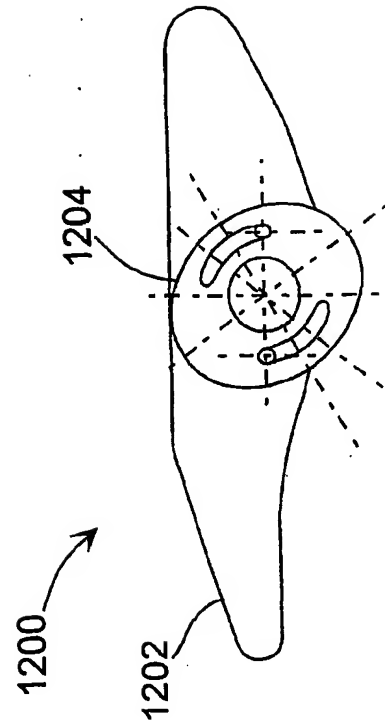


Fig. 124b

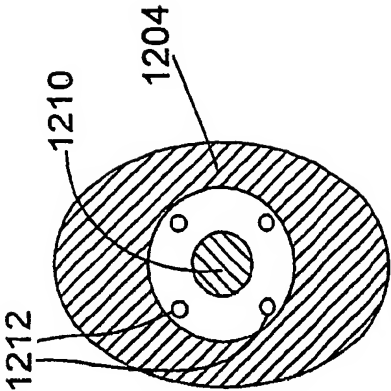


Fig. 127

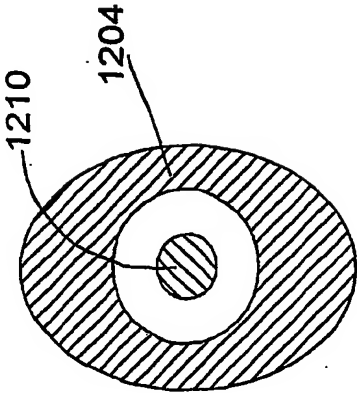


Fig. 126

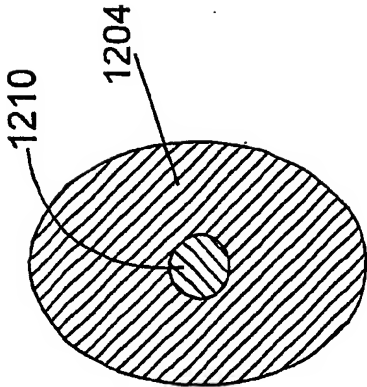


Fig. 125

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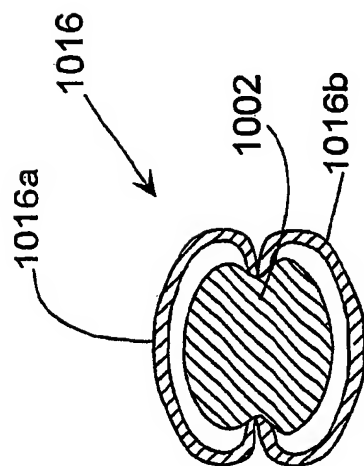


Fig. 128

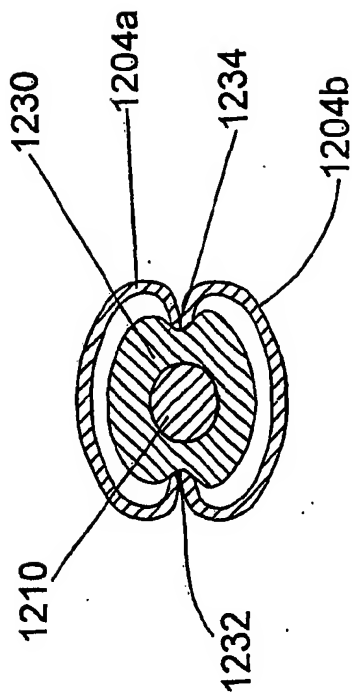


Fig. 129

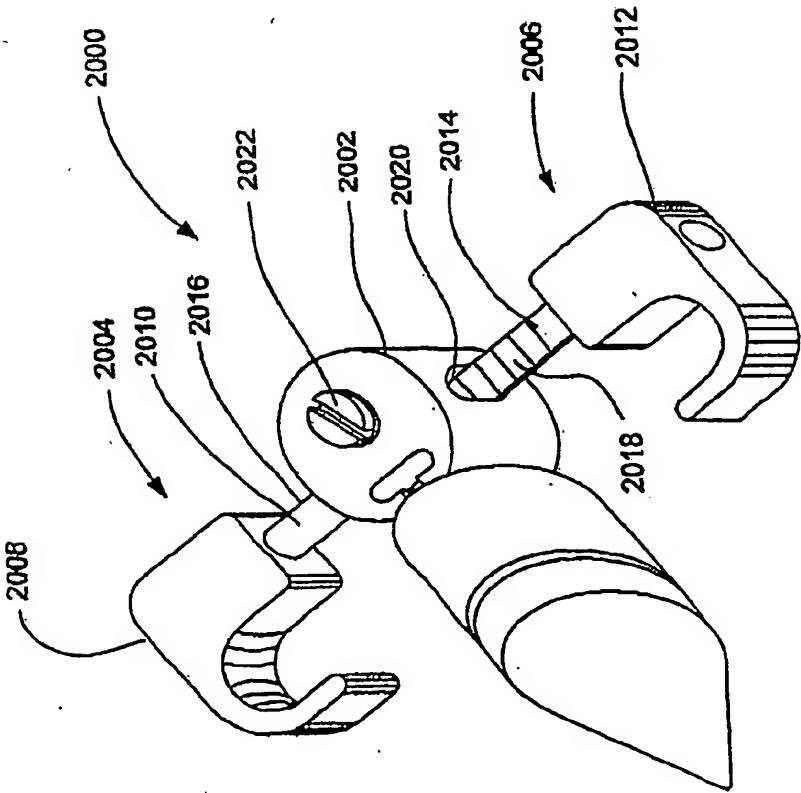


FIG. - 130

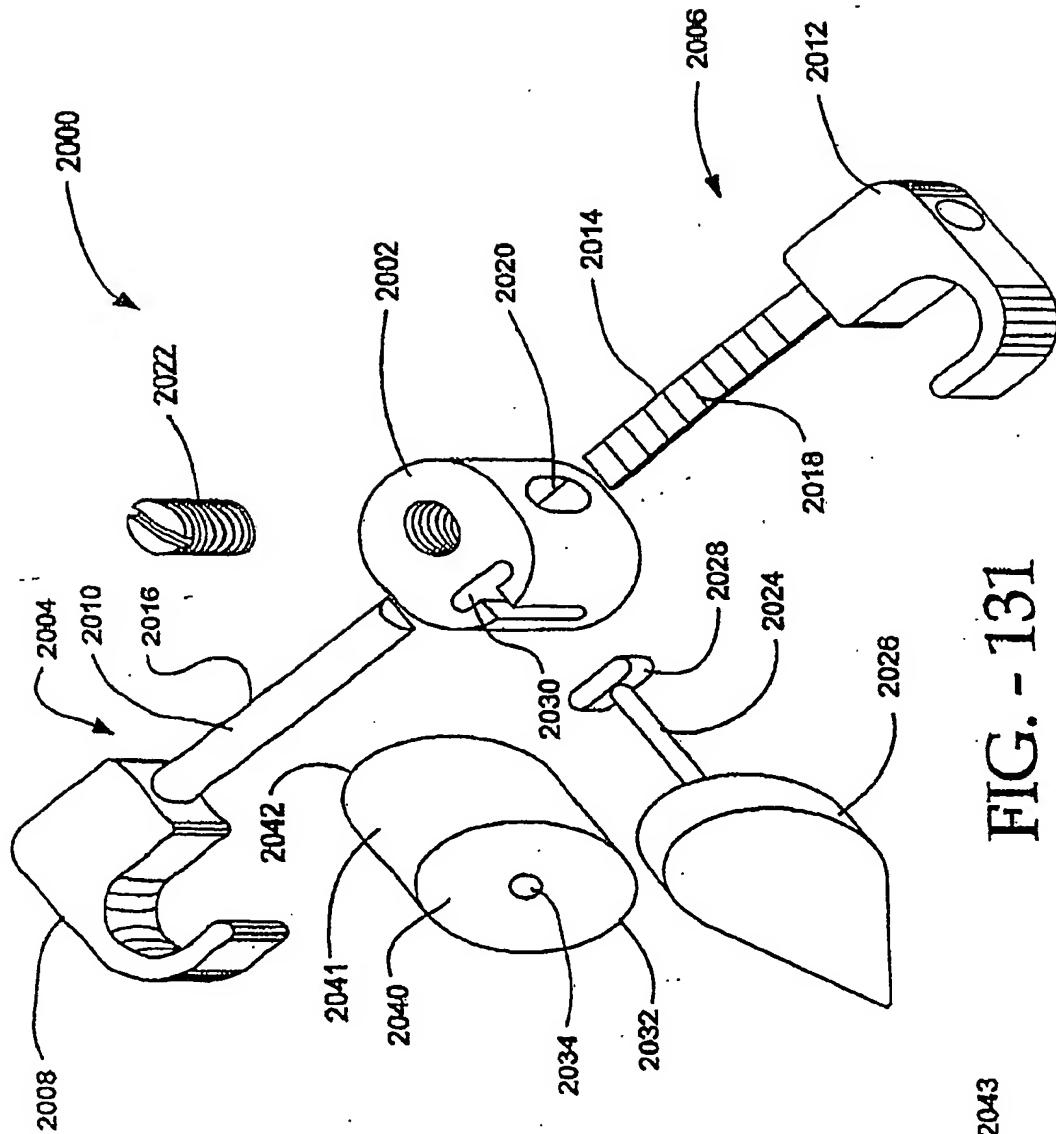


FIG. - 131

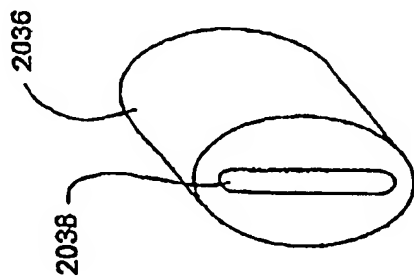


FIG. - 131a

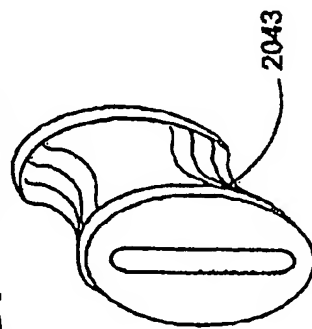


FIG. - 131b

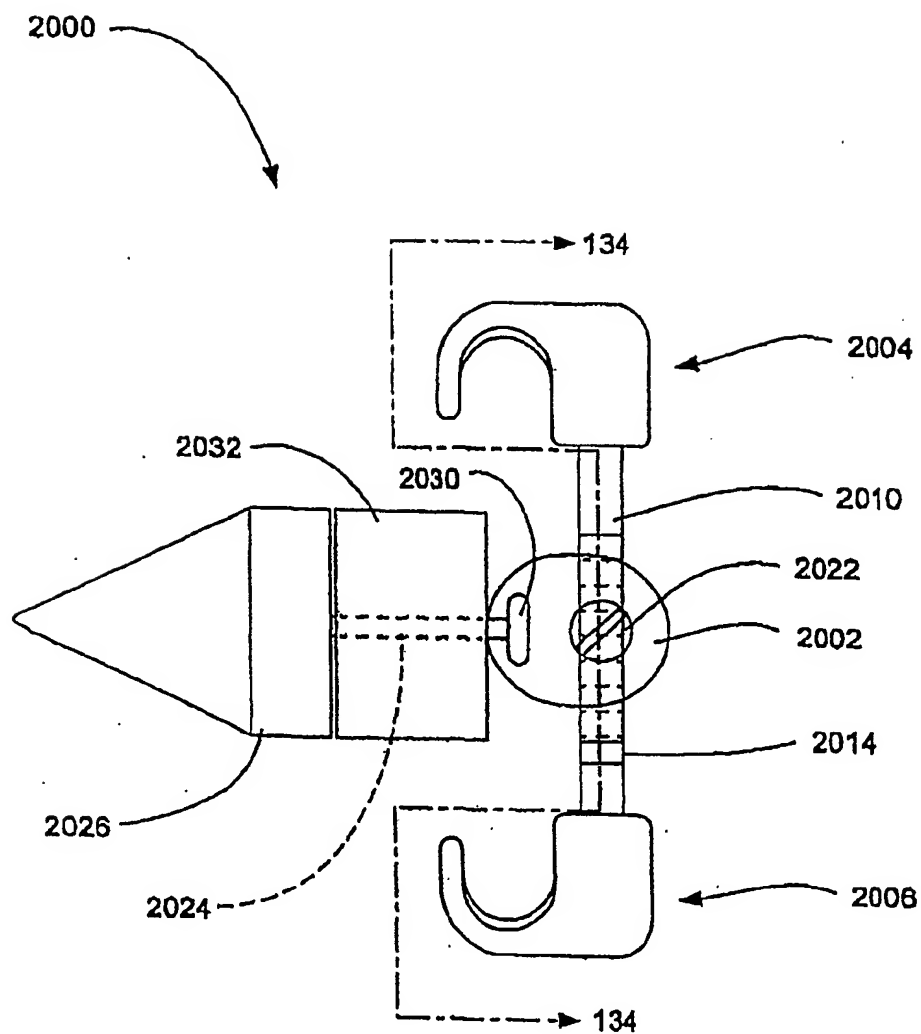


FIG. - 132

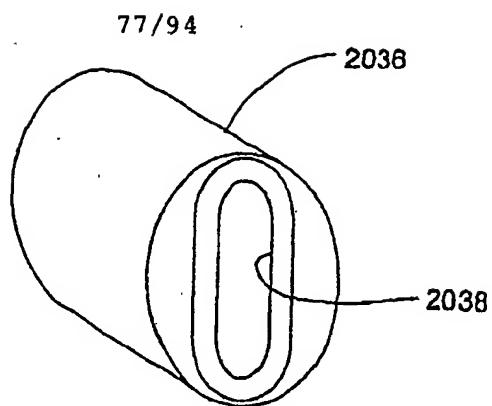


FIG. - 133a

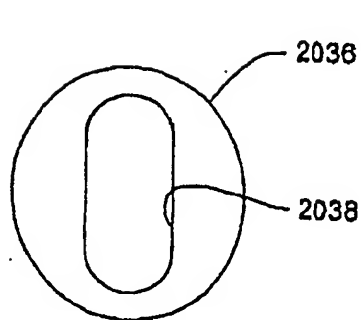


FIG. - 133b

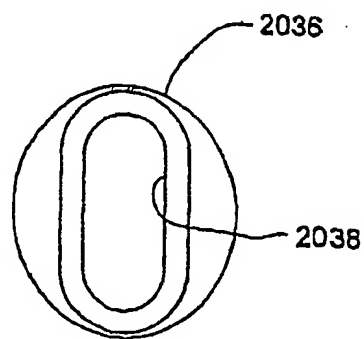


FIG. - 133c

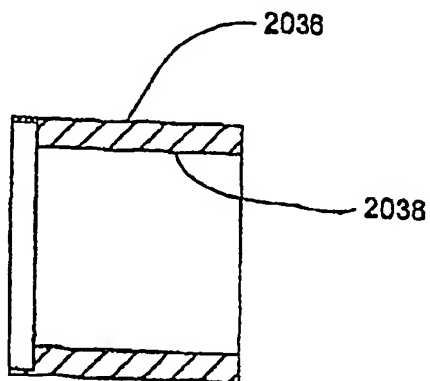


FIG. - 133d

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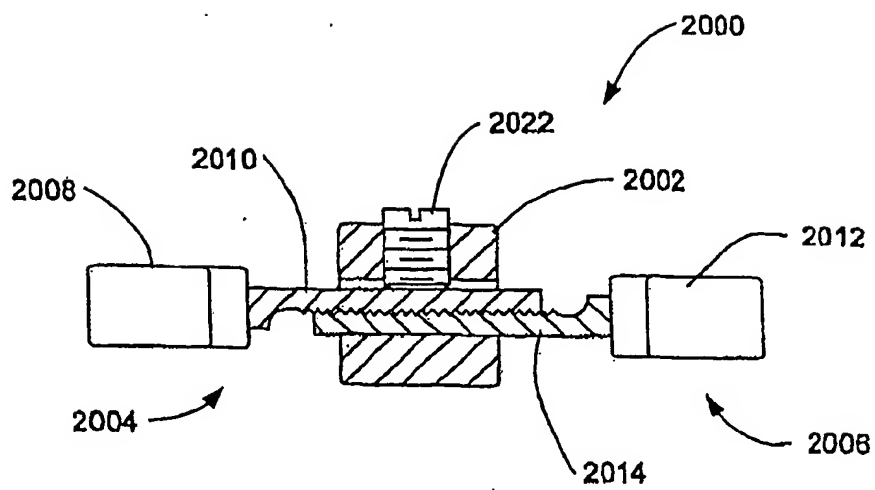


FIG. - 134

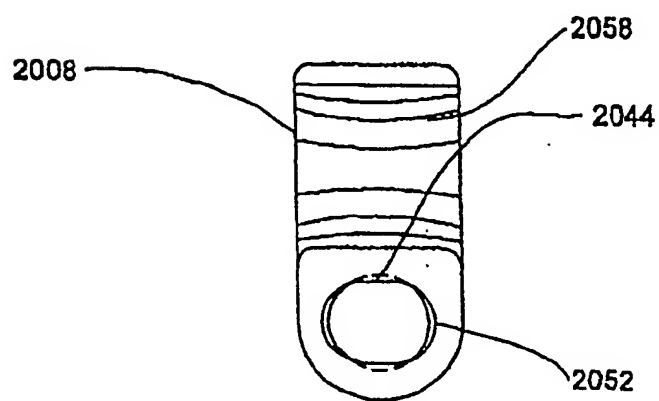


FIG. - 135a

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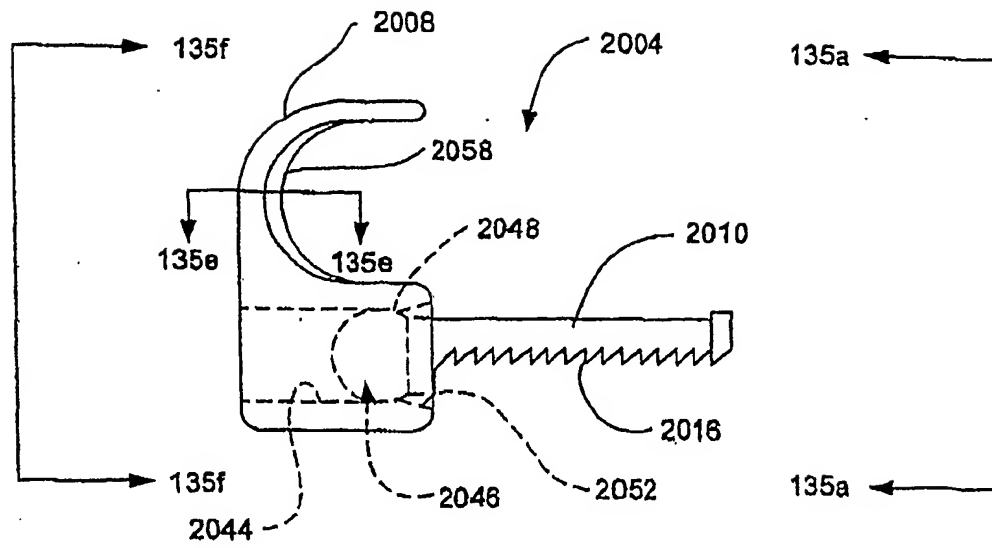


FIG. - 135b

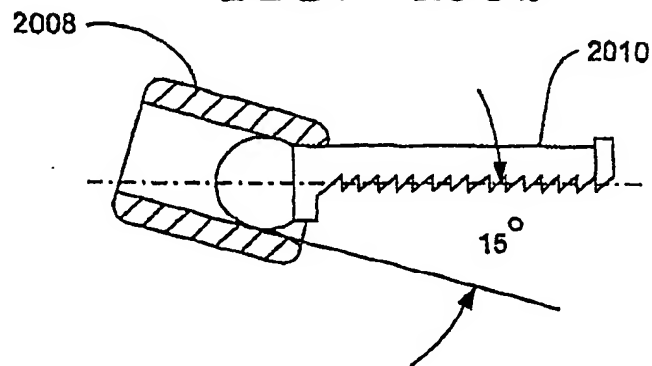


FIG. - 135c

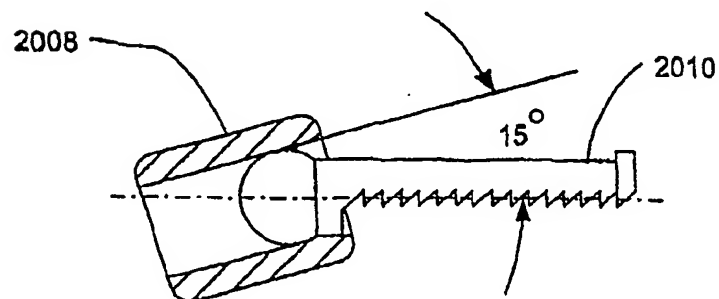


FIG. - 135d

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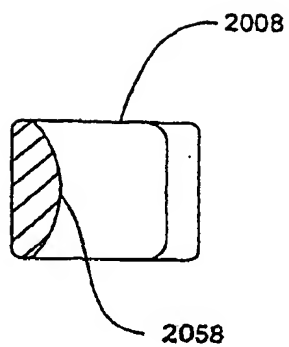


FIG. - 135e

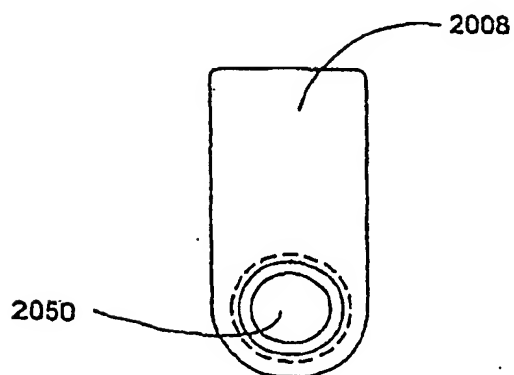


FIG. - 135f

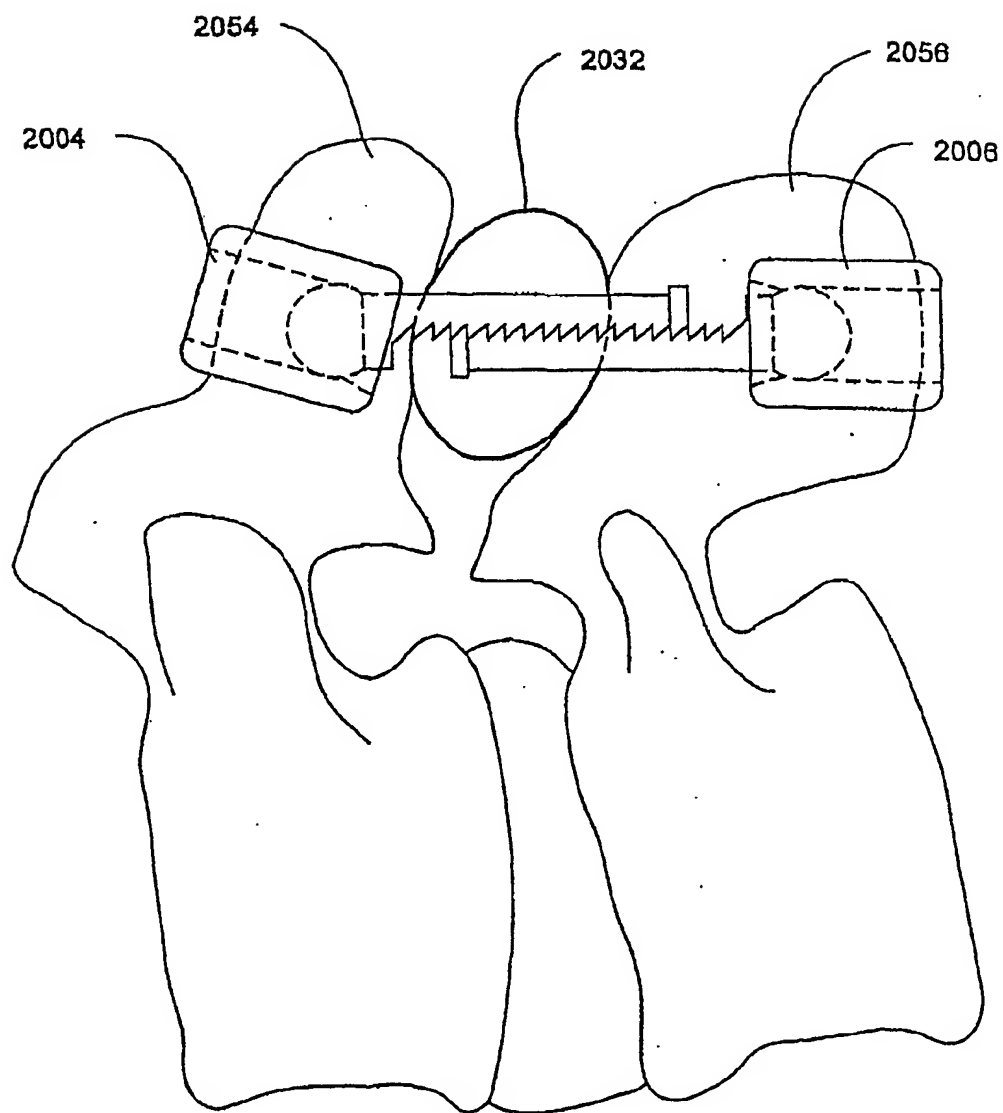


FIG. - 136

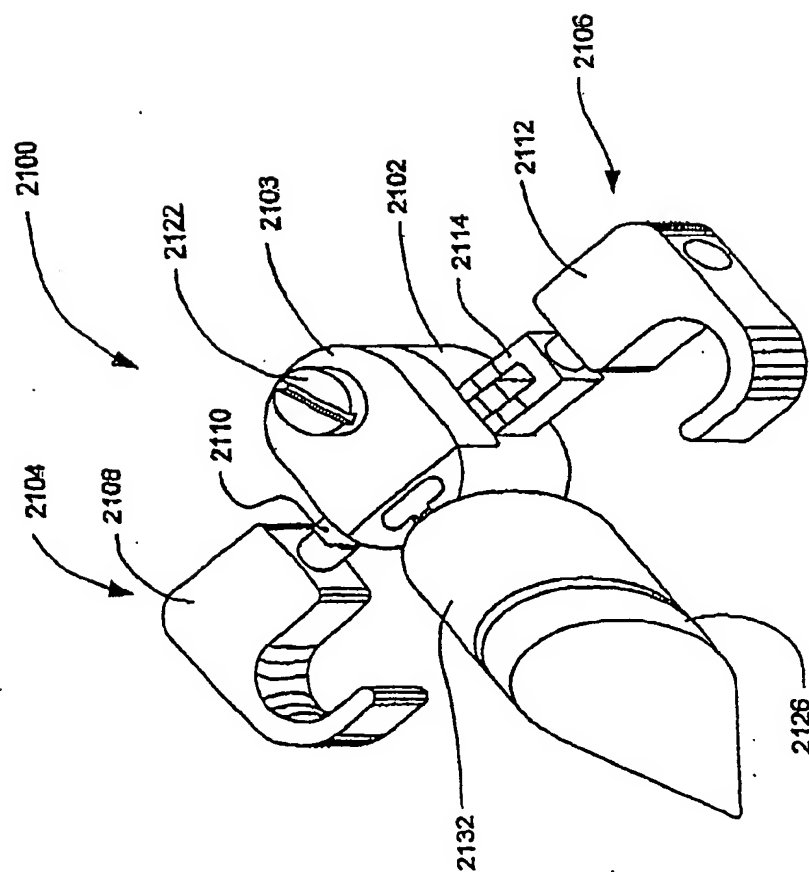
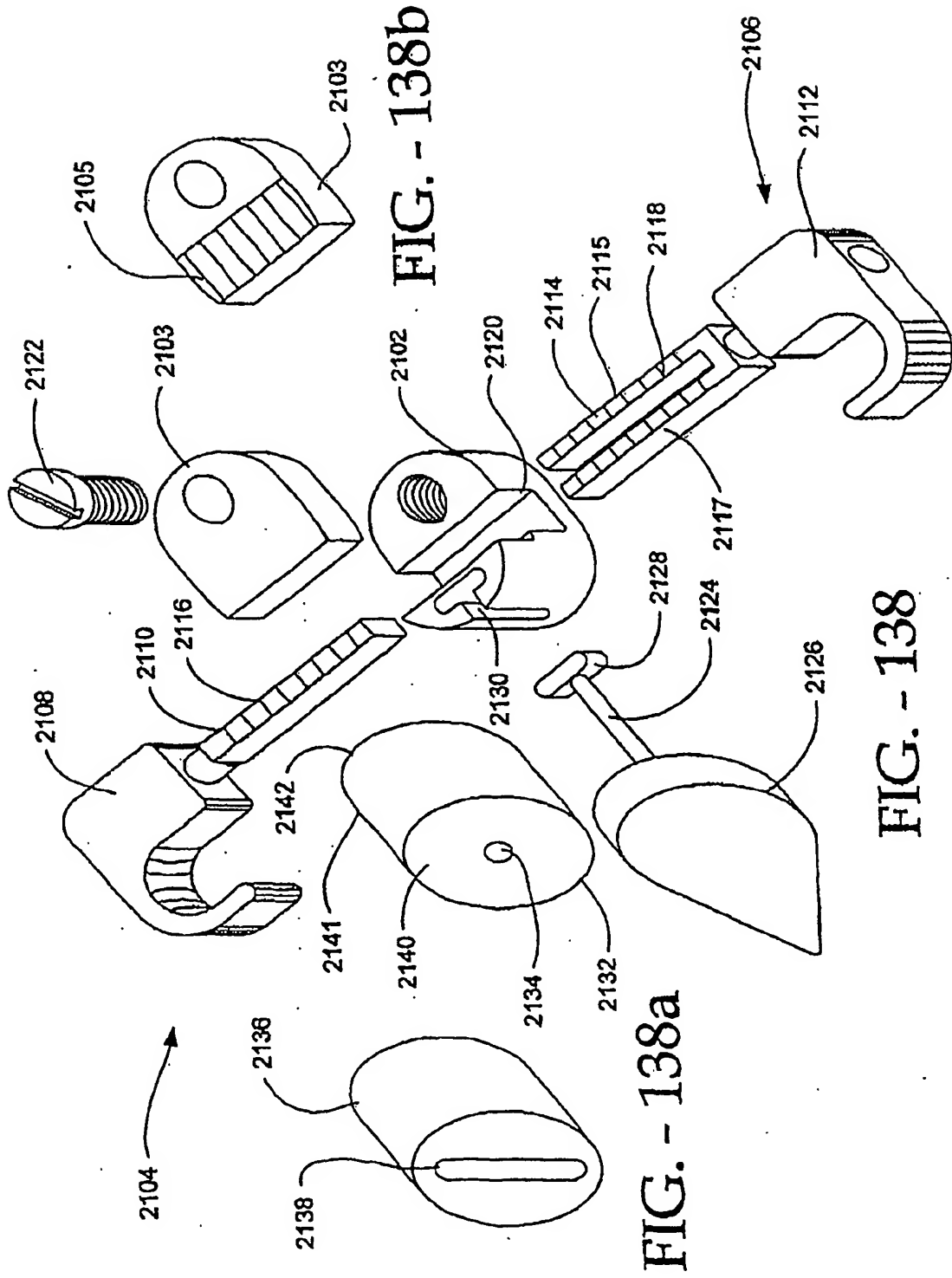


FIG. - 137



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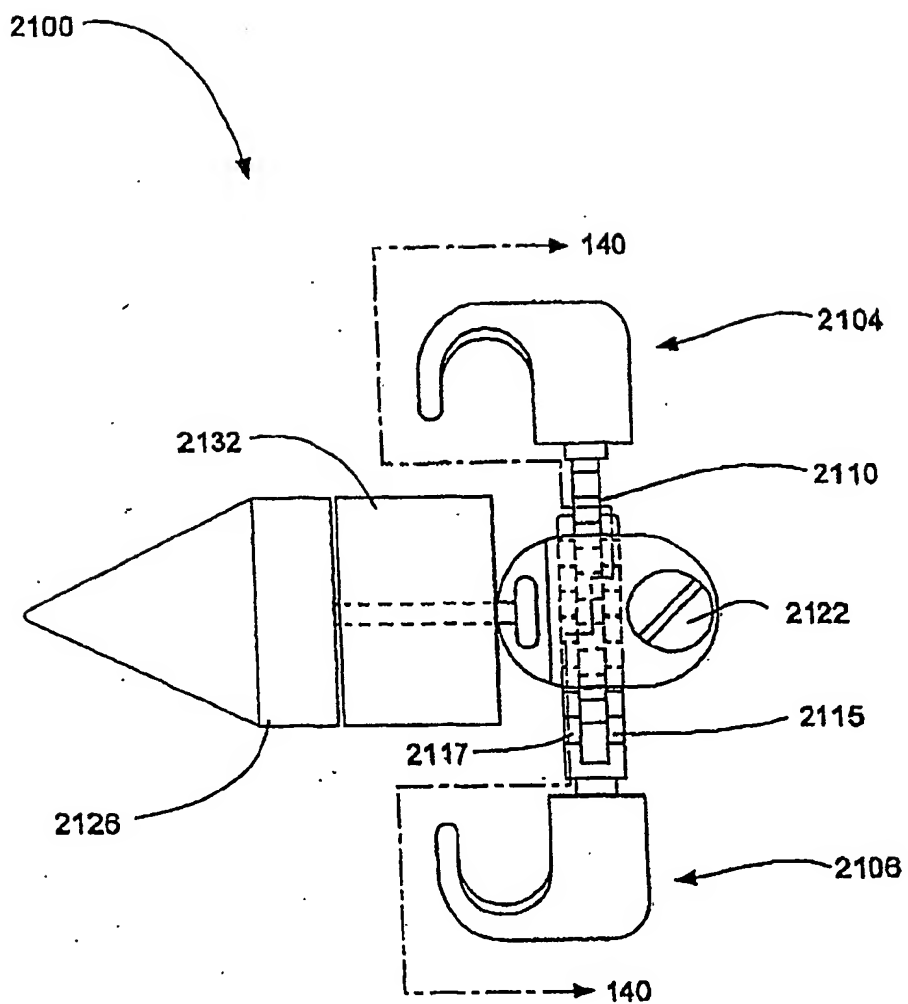


FIG. - 139

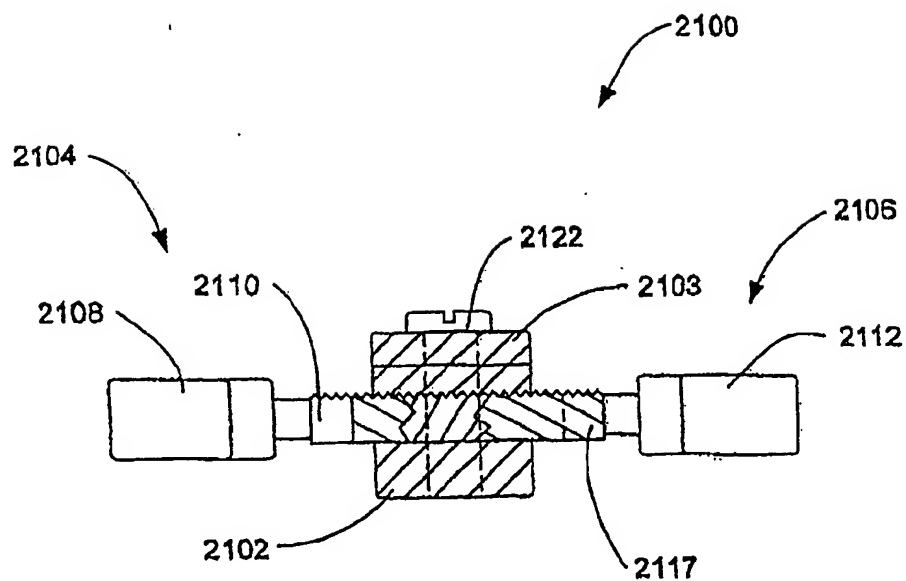


FIG. - 140

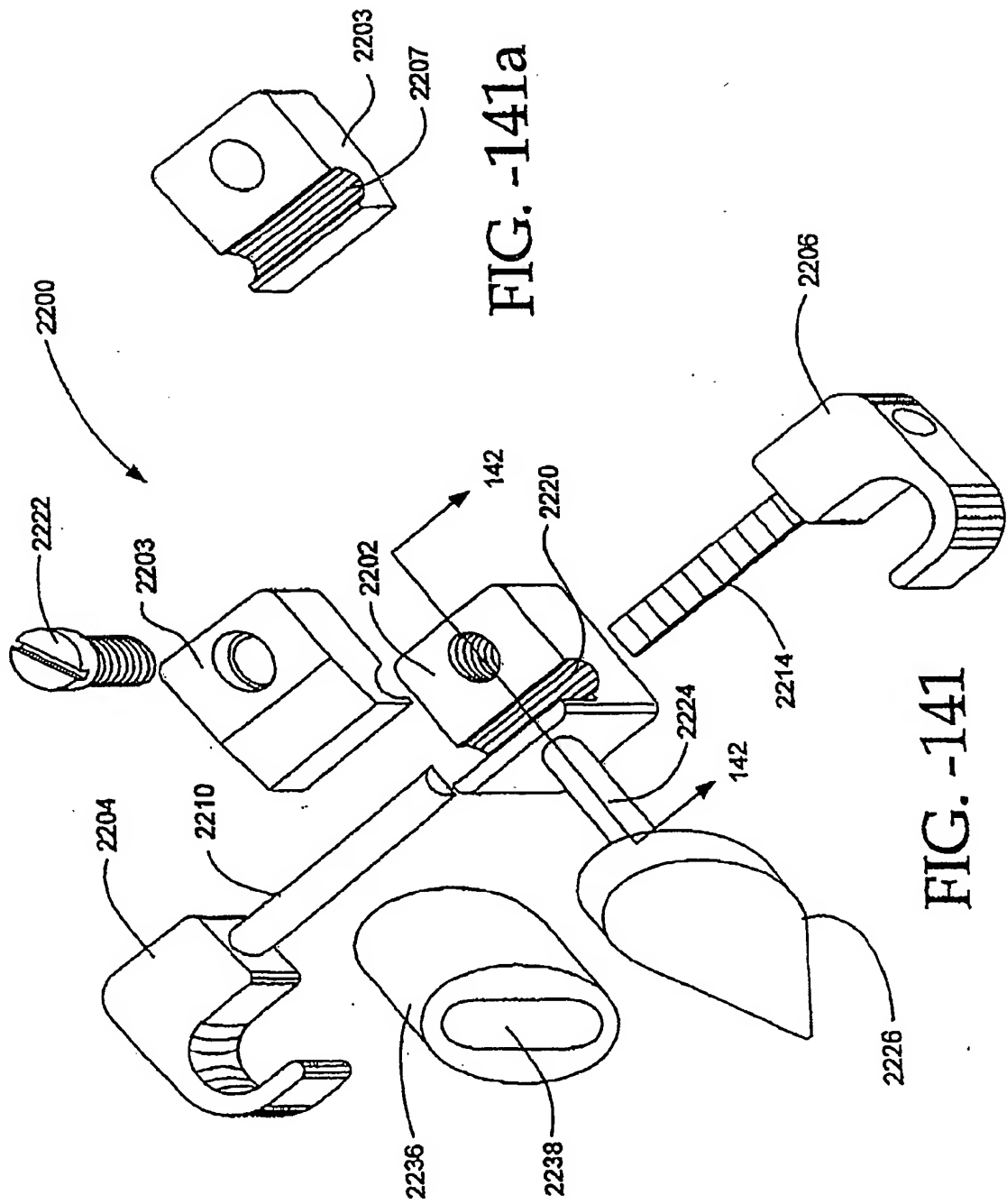


FIG. -141a

FIG. -141

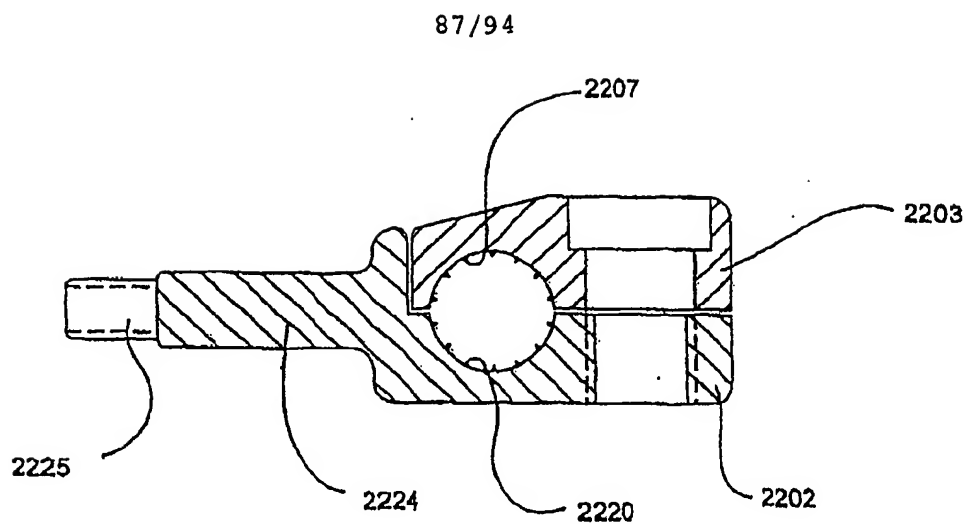


FIG. -142

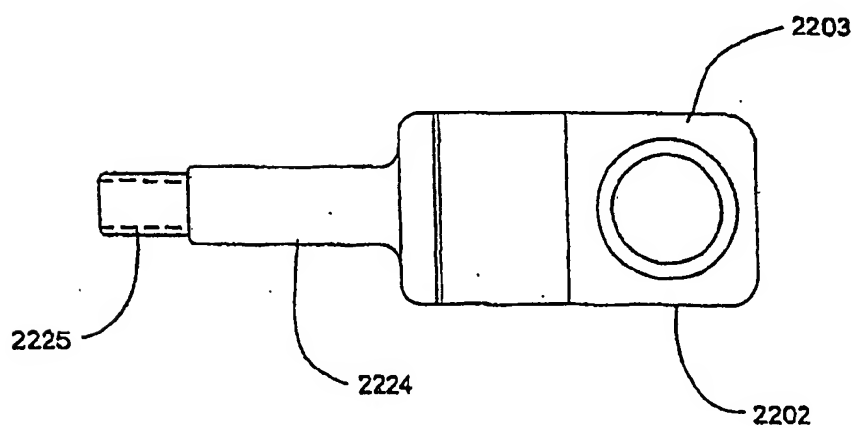


FIG. -143

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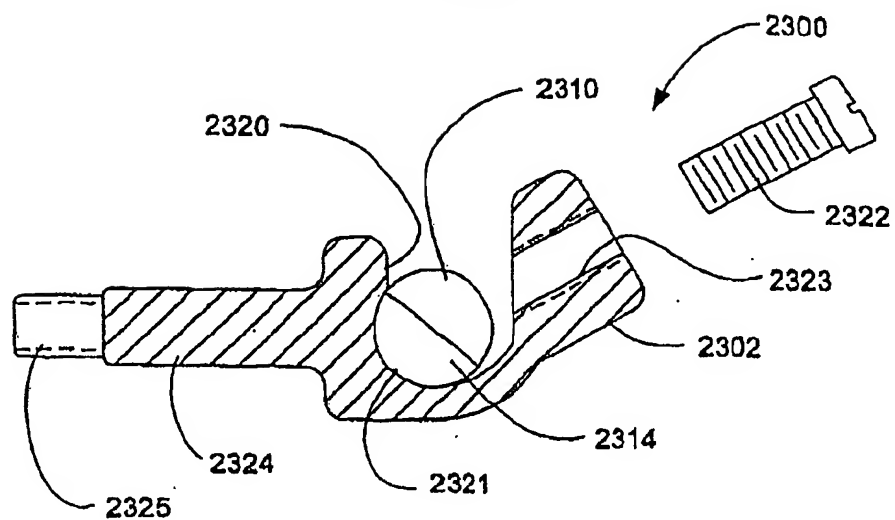


FIG. - 144

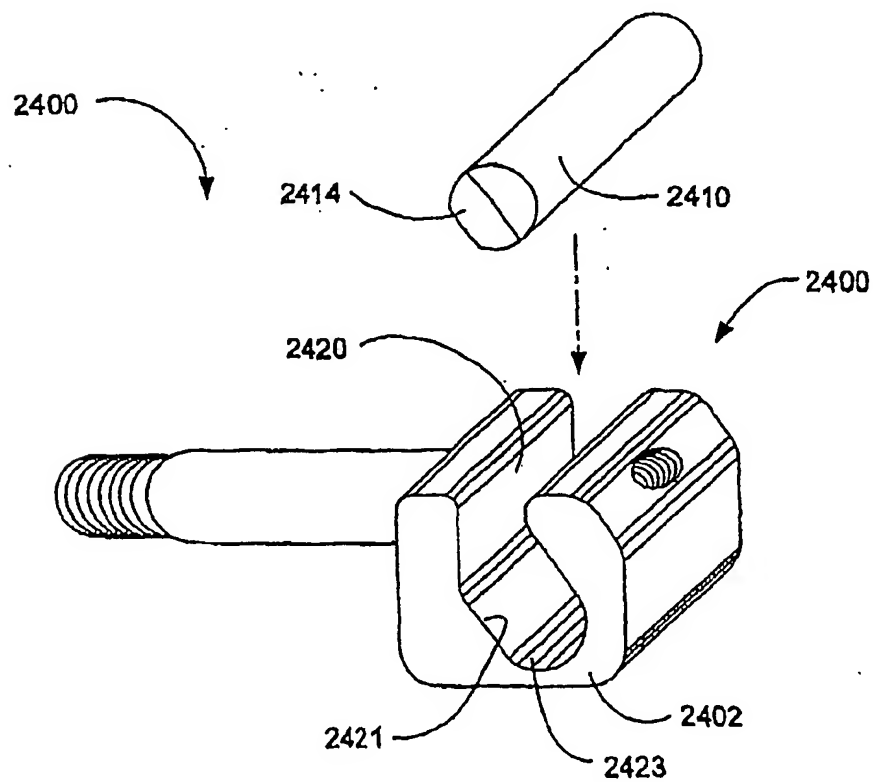
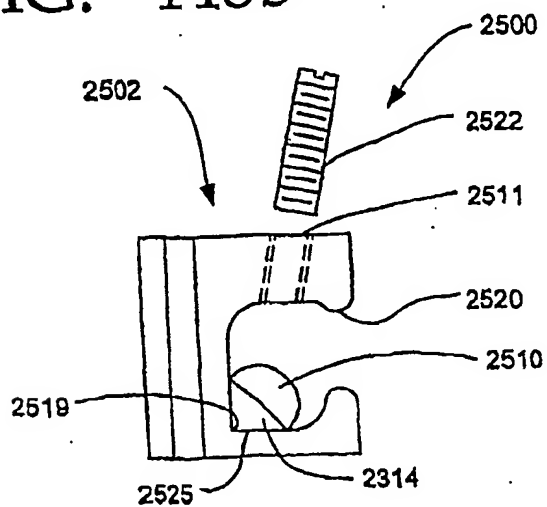
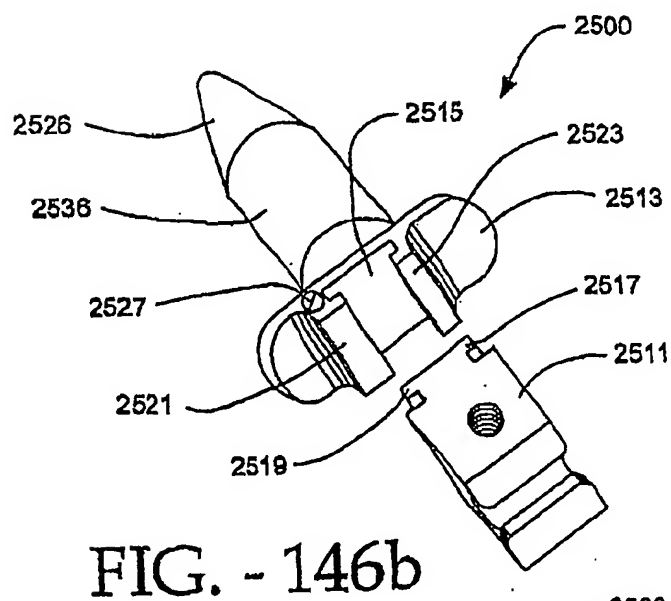
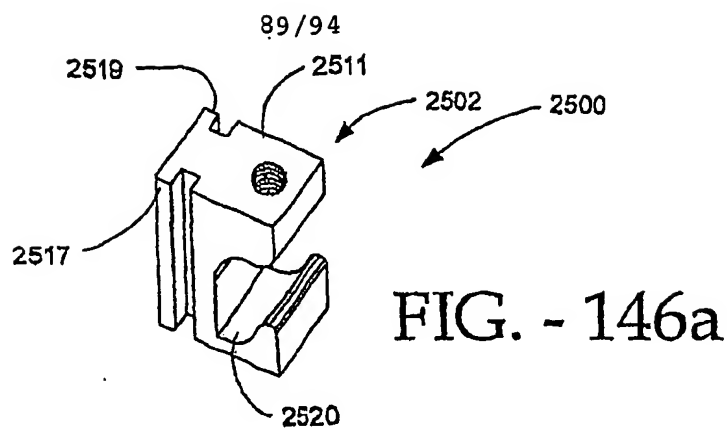


FIG. - 145



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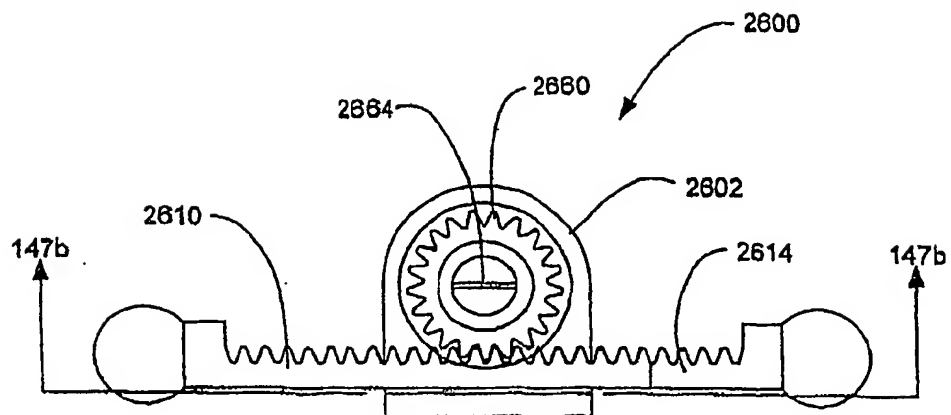


FIG. - 147a

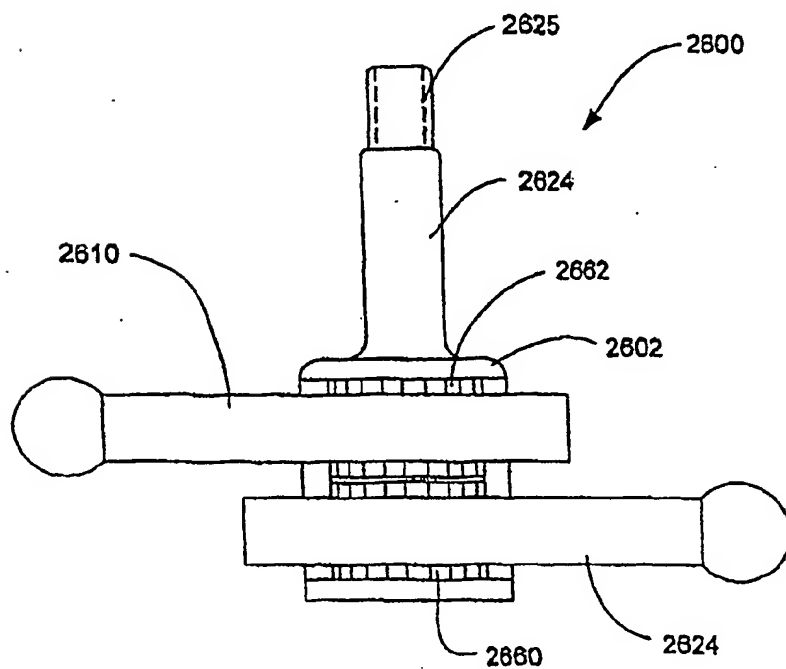


FIG. - 147b

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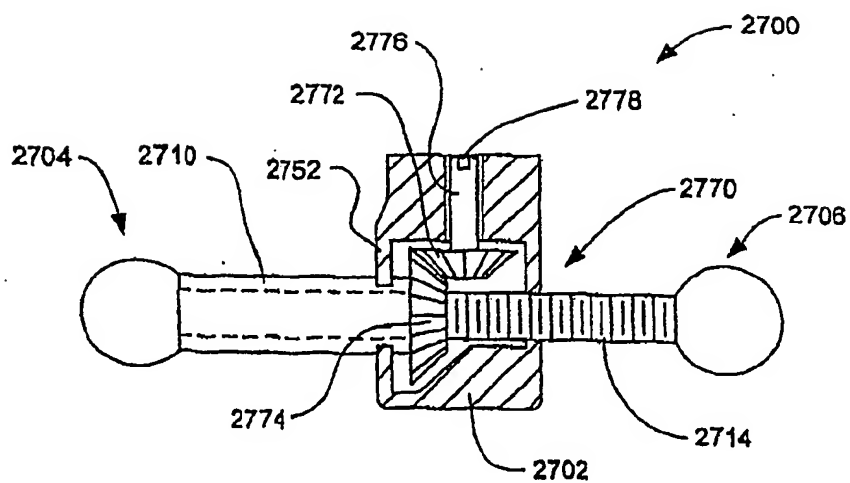


FIG. - 148a

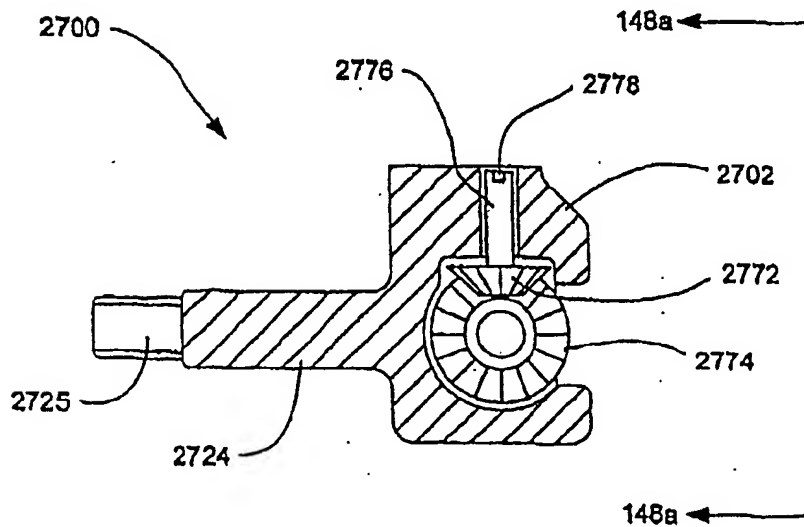
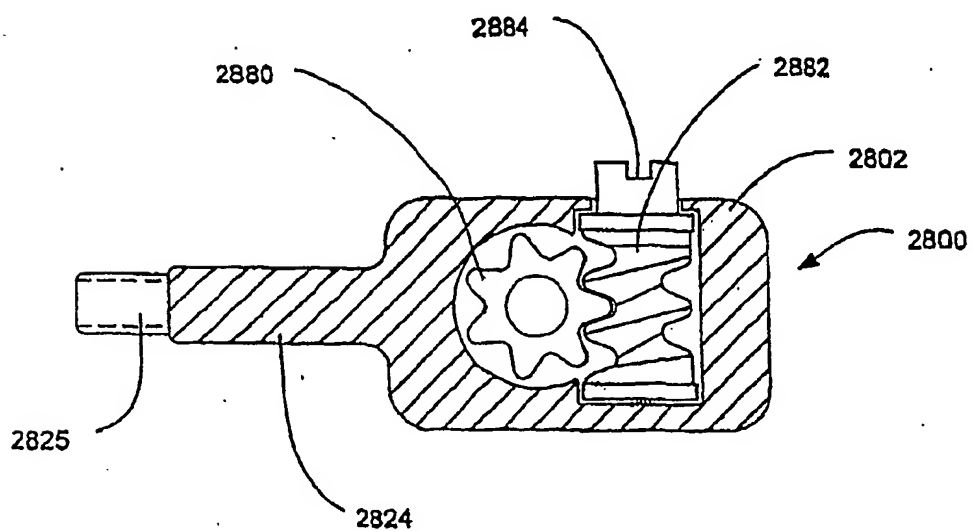
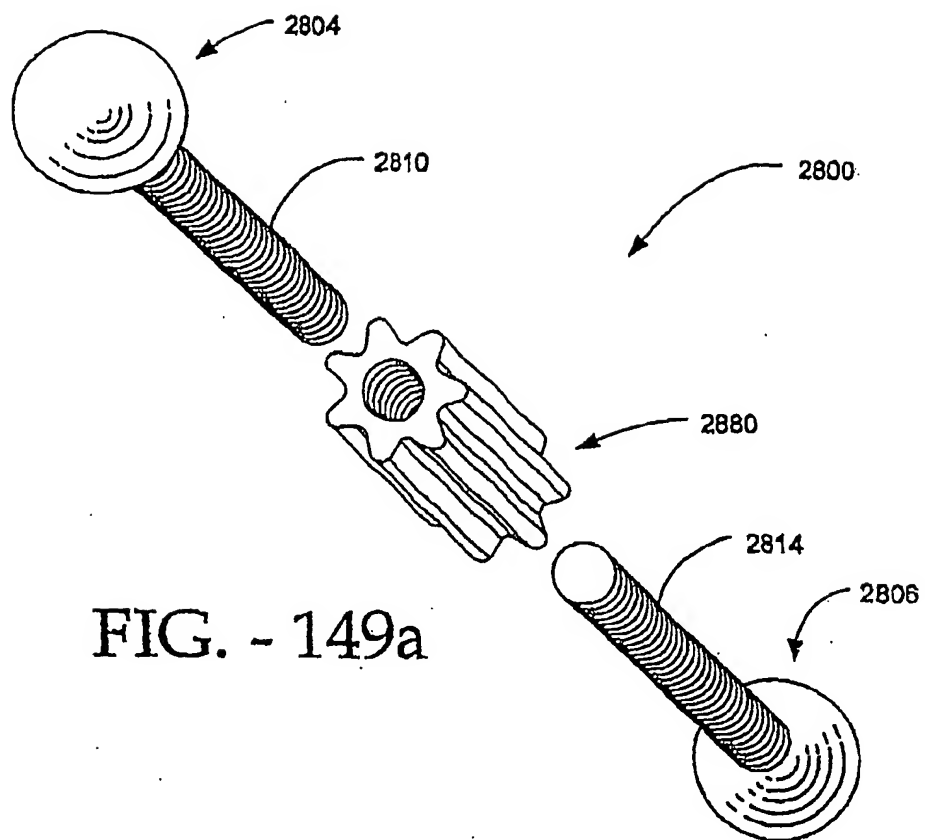
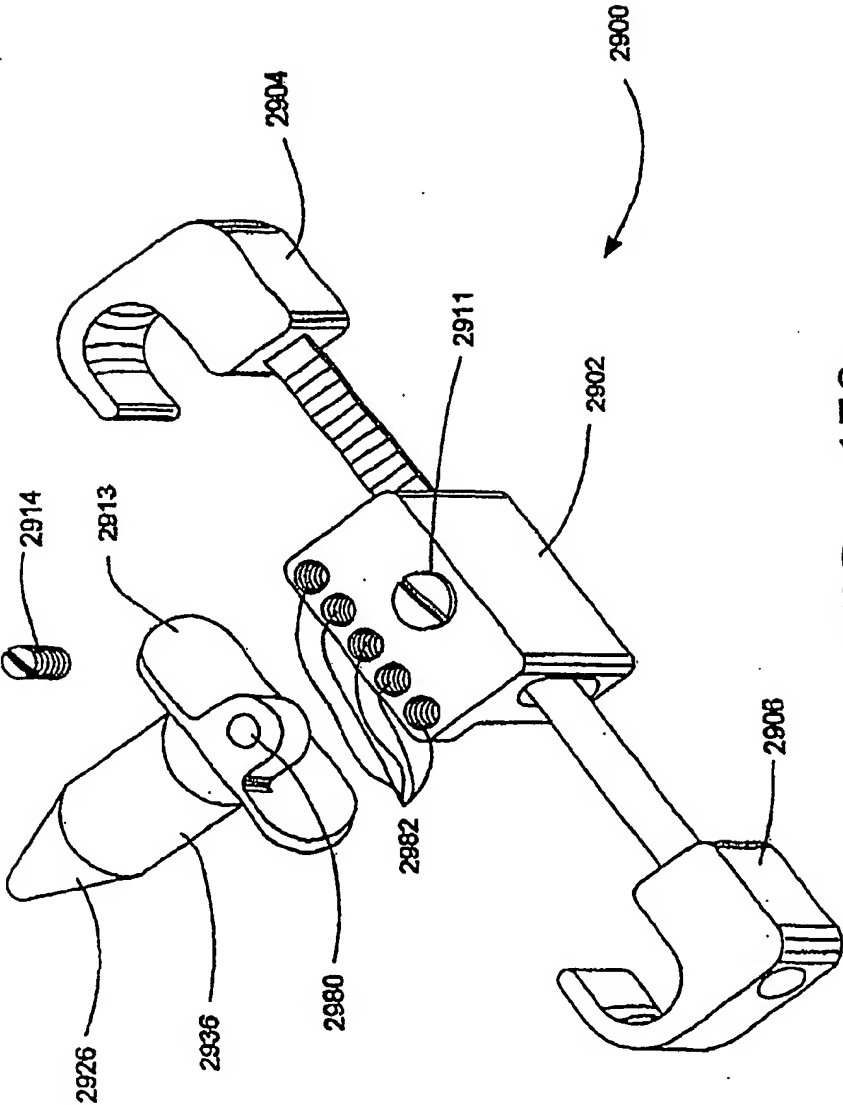


FIG. - 148b

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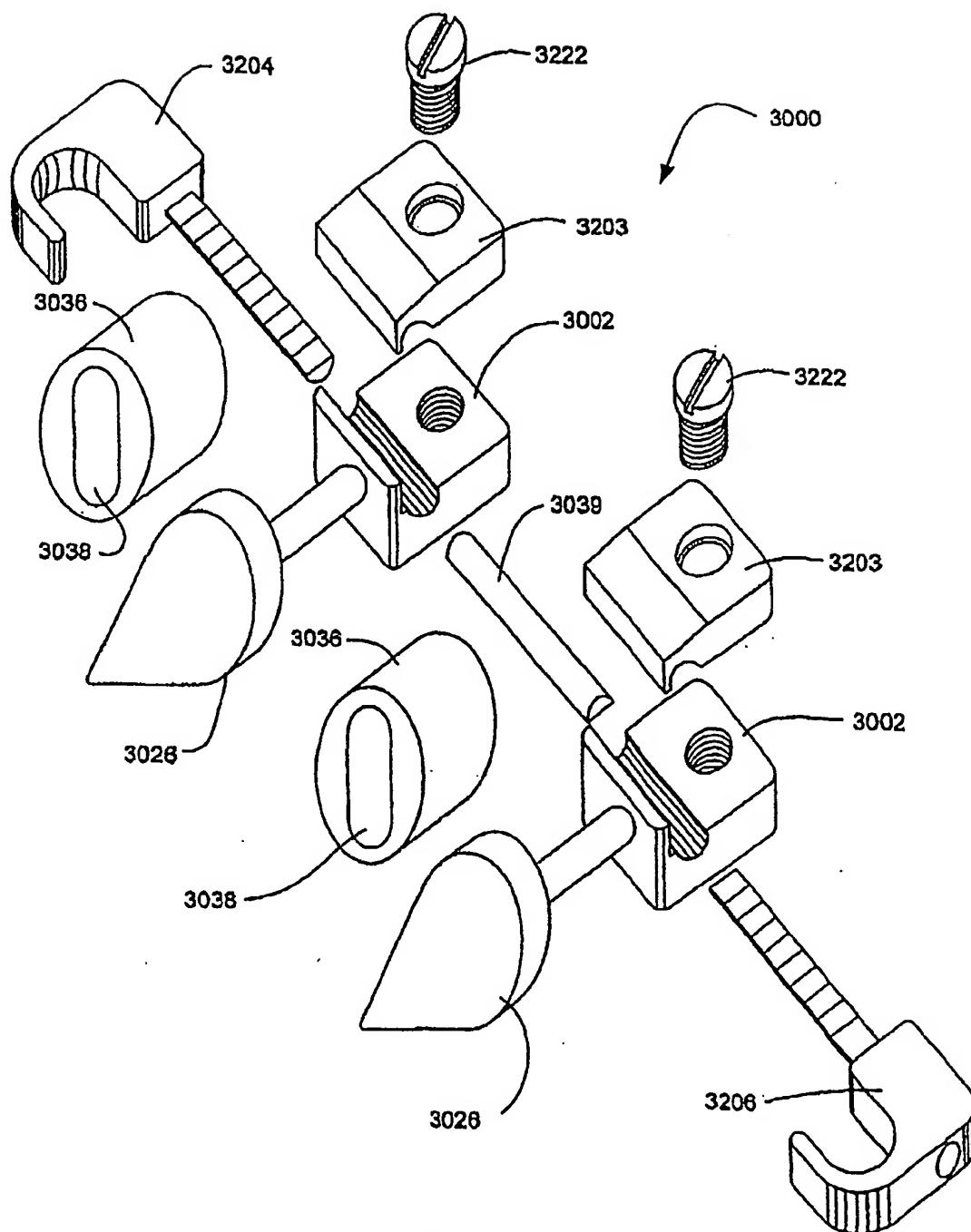


FIG. - 151

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US01/17295

| A. CLASSIFICATION OF SUBJECT MATTER | | |
|---|--|--|
| IPC(7) : A61B 17/70 US CL : 606/61 | | |
| According to International Patent Classification (IPC) or to both national classification and IPC | | |
| B. FIELDS SEARCHED | | |
| Minimum documentation searched (classification system followed by classification symbols) | | |
| U.S. : 606/60, 61, 71, 72 | | |
| Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched | | |
| Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) | | |
| C. DOCUMENTS CONSIDERED TO BE RELEVANT | | |
| Category* | Citation of document, with indication, where appropriate, of the relevant passages | Relevant to claim No. |
| X | US 5,645,599 A (SAMANI) 08 July 1997, Fig. 5. | 65-69 |
| X | US 5,496,318 A (HOWLAND et al) 05 March 1996, Fig. 7. | 65-67, 70 |
| X | US 2,677,369 A (KNOWLES) 04 May 1954, Figs. 3-4. | 65 |
| <input type="checkbox"/> Further documents are listed in the continuation of Box C. <input type="checkbox"/> See patent family annex. | | |
| * Special categories of cited documents: | "T" | later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention |
| "A" document defining the general state of the art which is not considered to be of particular relevance | "X" | document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone |
| "E" earlier document published on or after the international filing date | "Y" | document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art |
| "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) | "g" | document member of the same patent family |
| "O" document referring to an oral disclosure, use, exhibition or other means | | |
| "P" document published prior to the international filing date but later than the priority date claimed | | |
| Date of the actual completion of the international search | Date of mailing of the international search report | |
| 08 AUGUST 2001 | 31 AUG 2001 | |
| Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231 | Authorized officer | |
| Facsimile No. (703) 305-3230 | DAVID O. REIP <i>Diane Smith f</i> | |
| | Telephone No. (703) 308-0858 | |